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(21) International Application Number: PCT/US00/03288 (22) International Filing Date: 9 February 2000 (09.02.00) (30) Priority Data: 60/120,047 12 February 1999 (12.02.99) US 1090 7 February 2000 (07.02.00) SY (71) Applicant (for all designated States except US): VERTEX PHARMACEUTICALS INCORPORATED [US/US]; 130 Waverly Street, Cambridge, MA 02139-4242 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): HALE, Michael, R. [US/US]; 42 Sunset Road, Bedford, MA 01730 (US). BAKER, Christopher, T. [US/US]; 30 Judith Lane #11, Waltham, MA 02452 (US). STAMMERS, Timothy, A. [CA/CA]; Apartment 220, 2301 Lake Shore Boulevard West, Etobicoke, Ontario M1V 1A7 (CA). SHERRILL, Ronald, G. [US/US]; 139 Brannigan Place, Cary, NC 27511 (US). SPALTENSTEIN, Andrew [US/US]; 4105 Brewster Drive, Raleigh, NC 27606 (US). FURFINE, Eric, S. [US/US]; 4133 Livingstone Place, Durham, NC 27707 (US). MALTAIS, François [US/US]; 289 Highland Avenue #301, Somerville, MA 02144 (US). ANDREWS, Clarence, Webster, III [US/US]; 2604 Glendale Avenue, Durham, NC		27704 (US). MILLER, John, F. [US/US]; 310 Continental Drive, Durham, NC 27712 (US). SAMANO, Vicente [MX/US]; 410 Englewood Drive, Chapel Hill, NC 27514 (US). (74) Agents: HALEY, James, F., Jr.; Fish & Neave, 1251 Avenue of the Americas, New York, NY 10020 (US) et al. (81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>Without international search report and to be republished upon receipt of that report.</i>
(54) Title: INHIBITORS OF ASPARTYL PROTEASE (57) Abstract The present invention relates to a novel class of sulfonamides which are aspartyl protease inhibitors. In one embodiment, this invention relates to a novel class of HIV aspartyl protease inhibitors characterized by specific structural and physicochemical features. This invention also relates to pharmaceutical compositions comprising these compounds. The compounds and pharmaceutical compositions of this invention are particularly well suited for inhibiting HIV-1 and HIV-2 protease activity and consequently may be advantageously used as anti-viral agents against the HIV-1 and HIV-2 viruses. This invention also relates to methods for inhibiting the activity of HIV aspartyl protease using the compounds of this invention and methods for screening compounds for anti-HIV activity.		

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INHIBITORS OF ASPARTYL PROTEASE

TECHNICAL FIELD OF THE INVENTION

The present invention relates to a novel class of sulfonamides which are aspartyl protease inhibitors. In one embodiment, this invention relates to a novel class of HIV aspartyl protease inhibitors characterized by specific structural and physicochemical features. This invention also relates to pharmaceutical compositions comprising these compounds. The compounds and pharmaceutical compositions of this invention are particularly well suited for inhibiting HIV-1 and HIV-2 protease activity and consequently, may be advantageously used as anti-viral agents against the HIV-1 and HIV-2 viruses. This invention also relates to methods for inhibiting the activity of HIV aspartyl protease using the compounds of this invention and methods for screening compounds for anti-HIV activity.

BACKGROUND OF THE INVENTION

The human immunodeficiency virus ("HIV") is the causative agent for acquired immunodeficiency syndrome ("AIDS") -- a disease characterized by the destruction of the immune system, particularly of CD4⁺ T-cells, with attendant susceptibility to opportunistic infections -- and its precursor AIDS-related complex ("ARC") -- a syndrome characterized by symptoms such as persistent generalized lymphadenopathy, fever and weight loss.

As in the case of several other retroviruses, HIV encodes the production of a protease which carries

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out post-translational cleavage of precursor polypeptides in a process necessary for the formation of infectious virions (S. Crawford et al., "A Deletion Mutation in the 5' Part of the pol Gene of Moloney Murine Leukemia Virus Blocks Proteolytic Processing of the gag and pol Polyproteins", J. Virol., 53, p. 899 (1985)). These gene products include pol, which encodes the virion RNA-dependent DNA polymerase (reverse transcriptase), an endonuclease, HIV protease, and gag, which encodes the core-proteins of the virion (H. Toh et al., "Close Structural Resemblance Between Putative Polymerase of a Drosophila Transposable Genetic Element 17.6 and pol gene product of Moloney Murine Leukemia Virus", EMBO J., 4, p. 1267 (1985); L.H. Pearl et al., "A Structural Model for the Retroviral Proteases", Nature, pp. 329-351 (1987); M.D. Power et al., "Nucleotide Sequence of SRV-1, a Type D Simian Acquired Immune Deficiency Syndrome Retrovirus", Science, 231, p. 1567 (1986)).

A number of synthetic anti-viral agents have been designed to target various stages in the replication cycle of HIV. These agents include compounds which block viral binding to CD4⁺ T-lymphocytes (for example, soluble CD4), and compounds which interfere with viral replication by inhibiting viral reverse transcriptase (for example, didanosine and zidovudine (AZT)) and inhibit integration of viral DNA into cellular DNA (M.S. Hirsh and R.T. D'Aquila, "Therapy for Human Immunodeficiency Virus Infection", N.Eng.J.Med., 328, p. 1686 (1993)). However, such agents, which are directed primarily to early stages of viral replication, do not prevent the production of infectious virions in chronically infected cells. Furthermore, administration

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of some of these agents in effective amounts has led to cell-toxicity and unwanted side effects, such as anemia and bone marrow suppression.

More recently, the focus of anti-viral drug design has been to create compounds which inhibit the formation of infectious virions by interfering with the processing of viral polyprotein precursors. Processing of these precursor proteins requires the action of virus-encoded proteases which are essential for replication (Kohl, N.E. et al. "Active HIV Protease is Required for Viral Infectivity" Proc. Natl. Acad. Sci. USA, 85, p. 4686 (1988)). The anti-viral potential of HIV protease inhibition has been demonstrated using peptidyl inhibitors. Such peptidyl compounds, however, are typically large and complex molecules that tend to exhibit poor bioavailability and are not generally consistent with oral administration. Accordingly, the need still exists for compounds that can effectively inhibit the action of viral proteases, for use as agents for preventing and treating chronic and acute viral infections.

SUMMARY OF THE INVENTION

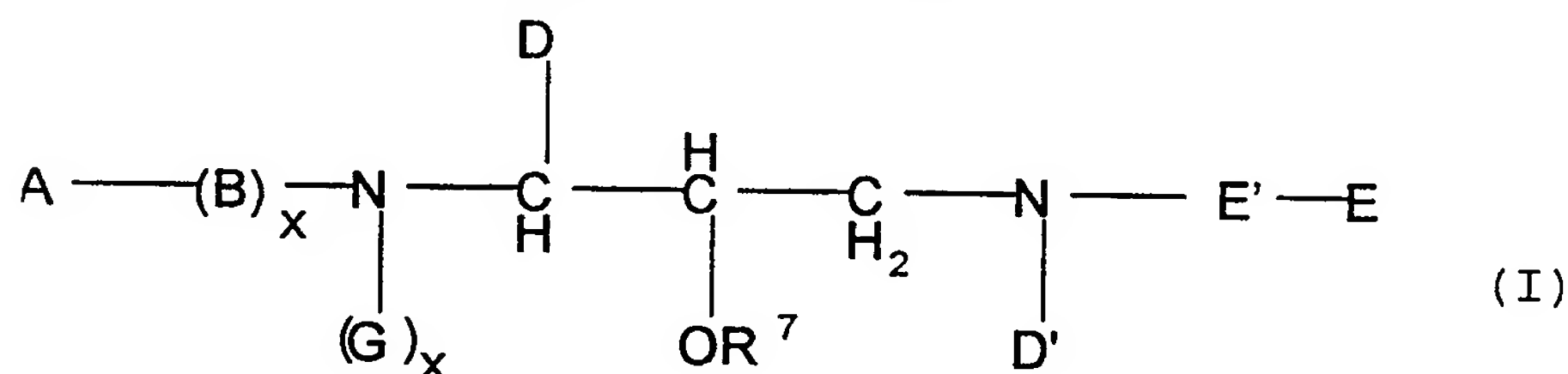
The present invention provides a novel class of compounds, and pharmaceutically acceptable derivatives thereof, that are useful as inhibitors of aspartyl proteases, in particular, HIV aspartyl protease. These compounds can be used alone or in combination with other therapeutic or prophylactic agents, such as anti-virals, antibiotics, immunomodulators or vaccines, for the treatment or prophylaxis of viral infection.

According to a preferred embodiment, the compounds of this invention are capable of inhibiting HIV

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viral replication in human CD₄⁺ T-cells. These compounds are useful as therapeutic and prophylactic agents to treat or prevent infection by HIV-1 and related viruses which may result in asymptomatic infection, AIDS-related complex ("ARC"), acquired immunodeficiency syndrome ("AIDS"), or similar disease of the immune system.

It is a principal object of this invention to provide a novel class of sulfonamides which are aspartyl protease inhibitors, and particularly, HIV aspartyl protease inhibitors. The novel sulfonamides of this invention are those of formula I:



wherein:

E' is -CO- or -SO₂-;

A is selected from H; Ht; -R¹-Ht; -R¹-C₁-C₆ alkyl, which is optionally substituted with one or more groups independently selected from hydroxy, C₁-C₄ alkoxy, Ht, -O-Ht, -NR²-CO-N(R²)₂; -SO₂-R² or -CO-N(R²)₂; -R¹-C₂-C₆ alkenyl, which is optionally substituted with one or more groups independently selected from hydroxy, C₁-C₄ alkoxy, Ht, -O-Ht, -NR²-CO-N(R²)₂ or -CO-N(R²)₂; or R⁷;

each R¹ is independently selected from -C(O)-, -S(O)₂-, -C(O)-C(O)-, -O-C(O)-, -O-S(O)₂-, -NR²-S(O)₂-, -NR²-C(O)- or -NR²-C(O)-C(O)-;

each Ht is independently selected from C₃-C₇ cycloalkyl; C₅-C₇ cycloalkenyl; C₆-C₁₄ aryl; or a 5-7 membered saturated or unsaturated heterocycle, containing one or more heteroatoms selected from N, O, or S; wherein

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said aryl or said heterocycle is optionally fused to Q;
 and wherein any member of said Ht is optionally
 substituted with one or more substituents independently
 selected from oxo, $-OR^2$, SR^2 , $-R^2$, $-N(R^2)(R^2)$, $-R^2-OH$, $-CN$,
 5 $-CO_2R^2$, $-C(O)-N(R^2)_2$, $-S(O)_2-N(R^2)_2$, $-N(R^2)-C(O)-R^2$, $-N(R^2)-$
 $C(O)O-R^2$, $-C(O)-R^2$, $-S(O)_n-R^2$, $-OCF_3$, $-S(O)_n-Q$,
 methylenedioxy, $-N(R^2)-S(O)_2(R^2)$, halo, $-CF_3$, $-NO_2$, Q, $-OQ$,
 $-OR^7$, $-SR^7$, $-R^7$, $-N(R^2)(R^7)$ or $-N(R^7)_2$;

each Q is independently selected from a 3-7 membered
 10 saturated, partially saturated or unsaturated carbocyclic
 ring system; or a 5-7 membered saturated, partially
 saturated or unsaturated heterocyclic ring containing one
 or more heteroatoms selected from O, N, or S; wherein Q
 is optionally substituted with one or more groups
 15 selected from oxo, $-OR^2$, $-R^2$, $-SO_2R^2$, $-SO_2-N(R^2)_2$, $-N(R^2)_2$,
 $-N(R^2)-C(O)-R^2$, $-R^2-OH$, $-CN$, $-CO_2R^2$, $-C(O)-N(R^2)_2$, halo,
 $-CF_3$;

each R^2 is independently selected from H, or C_1-C_4
 alkyl; and wherein said alkyl, when not a substituent of
 20 Q, is optionally substituted with Q or $-OR^3$; wherein when
 said R^2 is an $-OR^3$ substituted moiety, said R^3 in $-OR^3$ may
 not be $-OR^2$ substituted;

B, when present, is $-N(R^2)-C(R^3)_2-C(O)-$;

each x is independently 0 or 1;

25 each R^3 is independently selected from H, Ht, C_1-C_6
 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_3-C_6 cycloalkyl or C_5-
 C_6 cycloalkenyl; wherein any member of said R^3 , except H,
 is optionally substituted with one or more substituents
 selected from $-OR^2$, $-C(O)-NH-R^2$, $-S(O)_n-N(R^2)(R^2)$, $-N(R^2)_2$,
 30 $-N(R^2)-C(O)-O(R^2)$, $-N(R^2)-C(O)-N(R^2)$, $-N(R^2)-C(O)-(R^2)$, Ht,
 $-CN$, $-SR^2$, $-CO_2R^2$, or $NR^2-C(O)-R^2$;

each n is independently 1 or 2;

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G, when present, is selected from H, R⁷ or C₁-C₄ alkyl, or, when G is C₁-C₄ alkyl, G and R⁷ are optionally bound to one another either directly or through a C₁-C₃ linker to form a heterocyclic ring; or

5 when G is not present, the nitrogen to which G is attached is bound directly to the R⁷ group in -OR⁷ with the concomitant displacement of one -ZM group from R⁷;

D is selected from Q; C₁-C₆ alkyl optionally substituted with one or more groups selected from C₃-C₆ cycloalkyl, -OR², -S-Ht, -R³, -O-Q or Q; C₂-C₄ alkenyl optionally substituted with one or more groups selected from -OR², -S-Ht, -R³, -O-Q or Q; C₃-C₆ cycloalkyl optionally substituted with or fused to Q; or C₅-C₆ cycloalkenyl optionally substituted with or fused to Q;

15 D' is selected from C₁-C₁₅ alkyl, C₂-C₁₅ alkenyl or C₂-C₁₅ alkynyl, each of which contains one or more substituents selected from oxo, halo, -CF₃, -OCF₃, -NO₂, azido, -SH, -SR³, -N(R³)-N(R³)₂, -O-N(R³)₂, -(R³)N-O-(R³), -N(R³)₂, -CN, -CO₂R³, -C(O)-N(R³)₂, -S(O)_n-N(R³)₂, -N(R³)-C(O)-R³, -N(R³)-C(O)-N(R³)₂, -N(R³)-C(O)-S(R³), -C(O)-R³, -S(O)_n-R³, -N(R³)-S(O)_n(R³), -N(R³)-S(O)_n-N(R³)₂, -S-NR³-C(O)R³, -C(S)N(R³)₂, -C(S)R³, -NR³-C(O)OR³, -O-C(O)OR³, -O-C(O)N(R³)₂, -NR³-C(S)R³, =N-OH, =N-OR³, =N-N(R³)₂, =NR³, =NNR³C(O)N(R³)₂, , =NNR³C(O)OR³, =NNR³S(O)_n-N(R³)₂, -NR³-C(S)OR³, -NR³-C(S)N(R³)₂, -NR³-C[=N(R³)]-N(R³)₂, -N(R³)-C[=N-NO₂]-N(R³)₂, -N(R³)-C[=N-NO₂]-OR³, -N(R³)-C[=N-CN]-OR³, -N(R³)-C[=N-CN]-(R³)₂, -OC(O)R³, -OC(S)R³, -OC(O)N(R³)₂, -C(O)N(R³)-N(R³)₂, -O-C(O)N(R³)-N(R³)₂, -O-C(O)N(OR³)(R³), N(R³)-N(R³)C(O)R³, N(R³)-OC(O)R³, N(R³)-OC(O)R³, N(R³)-OC(O)R³, -OC(S)N(R³)₂, -OC(S)N(R³)(R³), or PO₃-R³; with the proviso that when R⁷ is H, E' is -SO₂-, G is H or alkyl, and when B is present or when B is not present and R¹ is

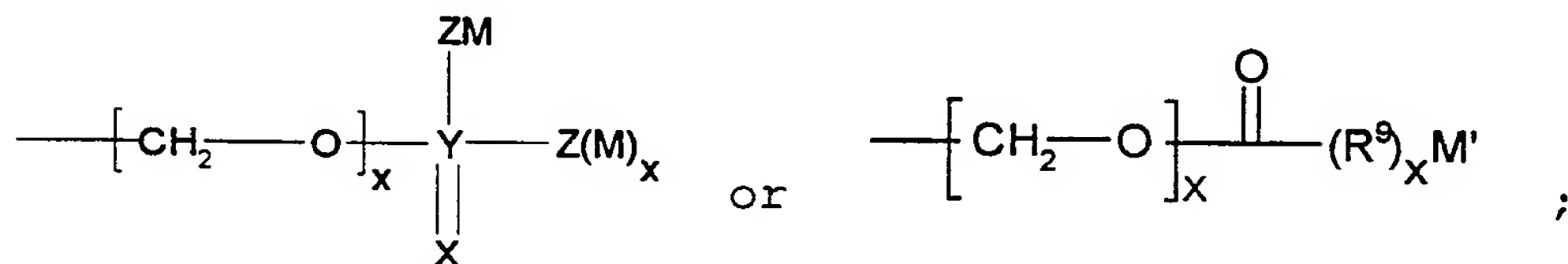
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-C(O)-, D' may not be C₁-C₁₅ alkyl substituted with one substituent selected from -N(R³)₂, -SR³ or -S(O)_n-R³, or substituted with two -N(R³)₂ substituents;

E is selected from Ht; O-Ht; Ht-Ht; Ht fused with
 5 Ht; -O-R³; -N(R²)(R³); C₁-C₆ alkyl optionally substituted with one or more groups selected from R⁴ or Ht; C₂-C₆ alkenyl optionally substituted with one or more groups selected from R⁴ or Ht; C₃-C₆ saturated carbocycle optionally substituted with one or more groups selected
 10 from R⁴ or Ht; or C₅-C₆ unsaturated carbocycle optionally substituted with one or more groups selected from R⁴ or Ht;

each R⁴ is independently selected from -OR², -OR³, -SR², -SOR², -SO₂R², -CO₂R², -C(O)-NHR², -C(O)-N(R²)₂,
 15 -C(O)-NR²(OR²), -S(O)₂-NHR², halo, -NR²-C(O)-R², -N(R²)₂ or -CN;

each R⁷ is independently selected from hydrogen,



20 wherein each M is independently selected from H, Li, Na, K, Mg, Ca, Ba, -N(R²)₄, C₁-C₁₂-alkyl, C₂-C₁₂-alkenyl, or -R⁶; wherein 1 to 4 -CH₂ radicals of the alkyl or alkenyl group, other than the -CH₂ that is bound to Z, is optionally replaced by a heteroatom group selected from
 25 O, S(O), S(O)₂, or N(R²); and wherein any hydrogen in said alkyl, alkenyl or R⁶ is optionally replaced with a substituent selected from oxo, -OR², -R², N(R²)₂, N(R²)₃, R²OH, -CN, -CO₂R², -C(O)-N(R²)₂, S(O)₂-N(R²)₂, N(R²)-C(O)-R²,

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$C(O)R^2$, $-S(O)_n-R^2$, OCF_3 , $-S(O)_n-R^6$, $N(R^2)-S(O)_2(R^2)$, halo, $-CF_3$, or $-NO_2$;

M' is H, C_1-C_{12} -alkyl, C_2-C_{12} -alkenyl, or $-R^6$; wherein 1 to 4 $-CH_2$ radicals of the alkyl or alkenyl group is optionally replaced by a heteroatom group selected from O, S, $S(O)$, $S(O)_2$, or $N(R^2)$; and wherein any hydrogen in said alkyl, alkenyl or R^6 is optionally replaced with a substituent selected from oxo, $-OR^2$, $-R^2$, $-N(R^2)_2$, $N(R^2)_3$, $-R^2OH$, $-CN$, $-CO_2R^2$, $-C(O)-N(R^2)_2$, $-S(O)_2-N(R^2)_2$, $-N(R^2)-C(O)-R^2$, $-C(O)R^2$, $-S(O)_n-R^2$, $-OCF_3$, $-S(O)_n-R^6$, $-N(R^2)-S(O)_2(R^2)$, halo, $-CF_3$, or $-NO_2$;

Z is O, S, $N(R^2)_2$, or, when M is not present, H.

Y is P or S;

X is O or S;

R^9 is $C(R^2)_2$, O or $N(R^2)$; and wherein when Y is S, Z is not S;

R^6 is a 5-6 membered saturated, partially saturated or unsaturated carbocyclic or heterocyclic ring system, or an 8-10 membered saturated, partially saturated or unsaturated bicyclic ring system; wherein any of said heterocyclic ring systems contains one or more heteroatoms selected from O, N, S, $S(O)_n$ or $N(R^2)$; and wherein any of said ring systems optionally contains 1 to 4 substituents independently selected from OH, C_1-C_4 alkyl, $-O-C_1-C_4$ alkyl or $-O-C(O)-C_1-C_4$ alkyl; and

each R^5 is independently selected from hydrogen, C_1-C_8 alkyl, C_2-C_8 alkenyl, C_2-C_8 alkynyl or Ht, wherein any R^5 , except for hydrogen, is optionally substituted with $-CF_3$, $-PO_3R^3$, azido or halo.

It is also an object of this invention to provide pharmaceutical compositions comprising the

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sulfonamides of formula (I) and methods for their use as inhibitors of HIV aspartyl protease.

DETAILED DESCRIPTION OF THE INVENTION

In order that the invention herein described
5 may be more fully understood, the following detailed description is set forth. In the description, the following terms are employed herein:

Unless expressly stated to the contrary, the terms "-SO₂-" and "-S(O)₂-" as used herein refer to a
10 sulfone or sulfone derivative (i.e., both appended groups linked to the S), and not a sulfinic ester.

For the compounds of formula I, and intermediates thereof, the stereochemistry of OR⁷ is defined relative to D on the adjacent carbon atom, when
15 the molecule is drawn in an extended zigzag representation (such as that drawn for compound of formula I). If both OR⁷ and D reside on the same side of the plane defined by the extended backbone of the compound, the stereochemistry of OR⁷ will be referred to
20 as "syn". If OR⁷ and D reside on opposite sides of that plane, the stereochemistry of OR⁷ will be referred to as "anti".

The term "alkyl", alone or in combination with any other term, refers to a straight-chain or branch-
25 chain saturated aliphatic hydrocarbon radical containing the specified number of carbon atoms, or where no number is specified, preferably from 1 to about 15 and more preferably from 1 to about 10 carbon atoms. Examples of alkyl radicals include, but are not limited to, methyl,
30 ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isoamyl, n-hexyl and the like.

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The term "alkenyl," alone or in combination with any other term, refers to a straight-chain or branched-chain mono- or poly-unsaturated aliphatic hydrocarbon radical containing the specified number of carbon atoms, or where no number is specified, preferably from 2 to about 18 carbon atoms and more preferably, from 2 to about 8 carbon atoms. Examples of alkenyl radicals include, but are not limited to, ethenyl, propenyl, isopropenyl, 1,4-butadienyl, pentenyl and the like.

The term "alkynyl," alone or in combination with any other term, refers to a straight-chain or branched-chain hydrocarbon radical having one or more triple bonds containing the specified number of carbon atoms, or where no number is specified, preferably from 2 to about 18 carbon atoms and more preferably, from 2 to about 8 carbon atoms. Examples of alkynyl radicals include, but are not limited to, ethynyl, propynyl, isopropynyl, butynyl, pentynyl and the like.

The term "alkoxy" refers to an alkyl ether radical, wherein the term "alkyl" is defined above. Examples of suitable alkyl ether radicals include, but are not limited to, methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, isobutoxy, sec-butoxy, tert-butoxy and the like.

The term "aryl," alone or in combination with any other term, refers to a carbocyclic aromatic radical (such as phenyl or naphthyl) containing the specified number of carbon atoms, preferably from 6-15 carbon atoms, and more preferably from 6-10 carbon atoms, optionally substituted with one or more substituents selected from C1-6 alkoxy, (for example methoxy), nitro, halogen, (for example chloro), amino, carboxylate and

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hydroxy. Examples of aryl radicals include, but are not limited to phenyl, naphthyl, indenyl, indanyl, azulenyl, fluorenyl, anthracenyl and the like.

The term "heterocyclyl" or "heterocycle" refers to a stable 3-7 membered monocyclic heterocyclic ring or 8-11 membered bicyclic heterocyclic ring which is either saturated or unsaturated, and which may be optionally benzofused if monocyclic. Each heterocycle consists of one or more carbon atoms and from one to four heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur. As used herein, the terms "nitrogen and sulfur heteroatoms" include any oxidized form of nitrogen and sulfur, and the quaternized form of any basic nitrogen. A heterocyclyl radical may be attached at any endocyclic carbon or heteroatom which results in the creation of a stable structure. Preferred heterocycles include 5-7 membered monocyclic heterocycles and 8-10 membered bicyclic heterocycles. Examples of such groups include imidazolyl, imidazolinoyl, imidazolidinyl, quinolyl, isoquinolyl, indolyl, indazolyl, indazolinolyl, perhydropyridazyl, pyridazyl, pyridyl, pyrrolyl, pyrrolinyl, pyrrolidinyl, pyrazolyl, pyrazinyl, quinoxolyl, piperidinyl, pyranyl, pyrazolinyl, piperazinyl, pyrimidinyl, pyridazinyl, morpholinyl, thiamorpholinyl, furyl, thienyl, triazolyl, thiazolyl, carbolinyl, tetrazolyl, thiazolidinyl, benzofuranoyl, thiamorpholinyl sulfone, oxazolyl, benzoxazolyl, oxopiperidinyl, oxopyrrolidinyl, oxoazepinyl, azepinyl, isoxozolyl, isothiazolyl, furazanyl, tetrahydropyranyl, tetrahydrofuranyl, thiazolyl, thiadiazoyl, dioxolyl, dioxinyl, oxathieryl, benzodioxolyl, dithieryl, thiophenyl, tetrahydrothiophenyl, sulfolanyl, dioxanyl,

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dioxolanyl, tetahydrofurodihydrofuranyl,
tetrahydropyranodihydrofuranyl, dihydropyranyl,
tetradydrofurofuranyl and tetrahydropyranofuranyl.

The term "pharmaceutically effective amount"
5 refers to an amount effective in treating a virus
infection, for example an HIV infection, in a patient
either as monotherapy or in combination with other
agents. The term "treating" as used herein refers to the
alleviation of symptoms of a particular disorder in a
10 patient or the improvement of an ascertainable
measurement associated with a particular disorder. The
term "prophylactically effective amount" refers to an
amount effective in preventing a virus infection, for
example an HIV infection, in a patient. As used herein,
15 the term "patient" refers to a mammal, including a human.

The terms "HIV protease" and "HIV aspartyl
protease" are used interchangeably and refer to the
aspartyl protease encoded by the human immunodeficiency
virus type 1 or 2. In a preferred embodiment of this
20 invention, these terms refer to the human
immunodeficiency virus type 1 aspartyl protease.

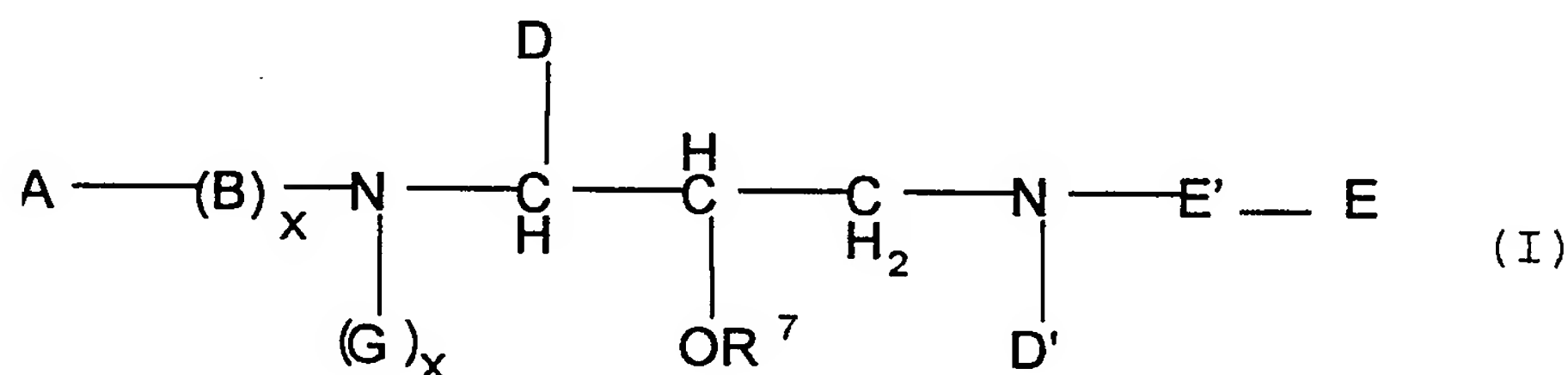
The term "thiocarbamates" refers to compounds
containing the functional group $\text{N-SO}_2\text{-O}$.

Combinations of substituents and variables
25 envisioned by this invention are only those that result
in the formation of stable compounds. The term "stable",
as used herein, refers to compounds which possess
stability sufficient to allow manufacture and
administration to a mammal by methods known in the art.
30 Typically, such compounds are stable at a temperature of
40°C or less, in the absence of moisture or other
chemically reactive conditions, for at least a week.

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This invention also envisions the quaternization of any basic nitrogen-containing groups of the compounds disclosed herein. The basic nitrogen can be quaternized with any agents known to those of ordinary skill in the art including, for example, lower alkyl halides, such as methyl, ethyl, propyl and butyl chloride, bromides and iodides; dialkyl sulfates including dimethyl, diethyl, dibutyl and diamyl sulfates; long chain halides such as decyl, lauryl, myristyl and stearyl chlorides, bromides and iodides; and aralkyl halides including benzyl and phenethyl bromides. Water or oil-soluble or dispersible products may be obtained by such quaternization.

The novel sulfonamides of this invention are those of formula I:



wherein:

E' is -CO- or -SO₂-;

A is selected from H; Ht; -R¹-Ht; -R¹-C₁-C₆ alkyl, which is optionally substituted with one or more groups independently selected from hydroxy, C₁-C₄ alkoxy, Ht, -O-Ht, -NR²-CO-N(R²)₂; -SO₂-R² or -CO-N(R²)₂; -R¹-C₂-C₆ alkenyl, which is optionally substituted with one or more groups independently selected from hydroxy, C₁-C₄ alkoxy, Ht, -O-Ht, -NR²-CO-N(R²)₂ or -CO-N(R²)₂; or R⁷;

each R¹ is independently selected from -C(O)-, -S(O)₂-, -C(O)-C(O)-, -O-C(O)-, -O-S(O)₂-, -NR²-S(O)₂-, -NR²-C(O)- or -NR²-C(O)-C(O)-;

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each Ht is independently selected from C₃-C₇ cycloalkyl; C₅-C₇ cycloalkenyl; C₆-C₁₄ aryl; or a 5-7 membered saturated or unsaturated heterocycle, containing one or more heteroatoms selected from N, O, or S; wherein
 5 said aryl or said heterocycle is optionally fused to Q; and wherein any member of said Ht is optionally substituted with one or more substituents independently selected from oxo, -OR², SR², -R², -N(R²)(R²), -R²-OH, -CN, -CO₂R², -C(O)-N(R²)₂, -S(O)₂-N(R²)₂, -N(R²)-C(O)-R², -N(R²)-
 10 C(O)O-R², -C(O)-R², -S(O)_n-R², -OCF₃, -S(O)_n-Q, methylenedioxy, -N(R²)-S(O)₂(R²), halo, -CF₃, -NO₂, Q, -OQ, -OR⁷, -SR⁷, -R⁷, -N(R²)(R⁷) or -N(R⁷)₂;

each Q is independently selected from a 3-7 membered saturated, partially saturated or unsaturated carbocyclic
 15 ring system; or a 5-7 membered saturated, partially saturated or unsaturated heterocyclic ring containing one or more heteroatoms selected from O, N, or S; wherein Q is optionally substituted with one or more groups selected from oxo, -OR², -R², -SO₂R², -SO₂-N(R²)₂, -N(R²)₂,
 20 -N(R²)-C(O)-R², -R²-OH, -CN, -CO₂R², -C(O)-N(R²)₂, halo, -CF₃;

each R² is independently selected from H, or C₁-C₄ alkyl,; and wherein said alkyl, when not a substituent of Q, is optionally substituted with Q or -OR³; wherein when
 25 said R² is an -OR³ substituted moiety, said R³ in -OR³ may not be -OR² substituted;

B, when present, is -N(R²)-C(R³)₂-C(O)-;

each x is independently 0 or 1;

each R³ is independently selected from H, Ht, C₁-C₆
 30 alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl or C₅-C₆ cycloalkenyl; wherein any member of said R³, except H, is optionally substituted with one or more substituents

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selected from $-OR^2$, $-C(O)-NH-R^2$, $-S(O)_n-N(R^2)(R^2)$, $-N(R^2)_2$, $-N(R^2)-C(O)-O(R^2)$, $-N(R^2)-C(O)-N(R^2)$, $-N(R^2)-C(O)-(R^2)$, Ht, $-CN$, $-SR^2$, $-CO_2R^2$, or $NR^2-C(O)-R^2$;

each n is independently 1 or 2;

5 G, when present, is selected from H, R^7 or C_1-C_4 alkyl, or, when G is C_1-C_4 alkyl, G and R^7 are optionally bound to one another either directly or through a C_1-C_3 linker to form a heterocyclic ring; or

10 when G is not present, the nitrogen to which G is attached is bound directly to the R^7 group in $-OR^7$ with the concomitant displacement of one $-ZM$ group from R^7 ;

D is selected from Q; C_1-C_6 alkyl optionally substituted with one or more groups selected from C_3-C_6 cycloalkyl, $-OR^2$, $-S-Ht$, $-R^3$, $-O-Q$ or Q; C_2-C_4 alkenyl optionally substituted with one or more groups selected from $-OR^2$, $-S-Ht$, $-R^3$, $-O-Q$ or Q; C_3-C_6 cycloalkyl optionally substituted with or fused to Q; or C_5-C_6 cycloalkenyl optionally substituted with or fused to Q;

20 D' is selected from C_1-C_{15} alkyl, C_2-C_{15} alkenyl or C_2-C_{15} alkynyl, each of which contains one or more substituents selected from oxo, halo, $-CF_3$, $-OCF_3$, $-NO_2$, azido, $-SH$, $-SR^3$, $-N(R^3)-N(R^3)_2$, $-O-N(R^3)_2$, $-(R^3)N-O-(R^3)$, $-N(R^3)_2$, $-CN$, $-CO_2R^3$, $-C(O)-N(R^3)_2$, $-S(O)_n-N(R^3)_2$, $-N(R^3)-C(O)-R^3$, $-N(R^3)-C(O)-N(R^3)_2$, $-N(R^3)-C(O)-S(R^3)$, $-C(O)-R^3$, $-S(O)_n-R^3$, $-N(R^3)-S(O)_n(R^3)$, $-N(R^3)-S(O)_n-N(R^3)_2$, $-S-NR^3-C(O)R^3$, $-C(S)N(R^3)_2$, $-C(S)R^3$, $-NR^3-C(O)OR^3$, $-O-C(O)OR^3$, $-O-C(O)N(R^3)_2$, $-NR^3-C(S)R^3$, $=N-OH$, $=N-OR^3$, $=N-N(R^3)_2$, $=NR^3$, $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$, $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-OR^3$, $-N(R^3)-C[=N-CN]-OR^3$, $-N(R^3)-C[=N-CN]-(R^3)_2$, $-OC(O)R^3$, $-OC(S)R^3$, $-OC(O)N(R^3)_2$, $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-N(R^3)_2$, $O-C(O)N(OR^3)(R^3)$,

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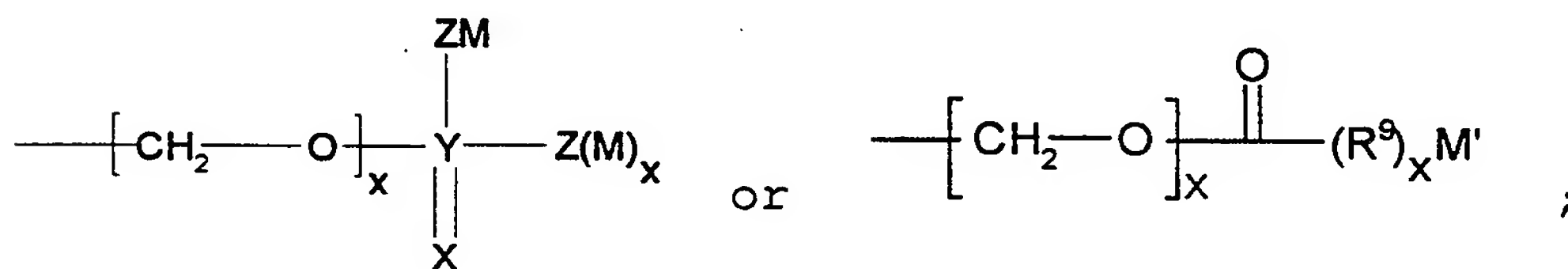
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$N(R^3)-N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $-OC(S)N(R^3)_2$, $-OC(S)N(R^3)(R^3)$, or PO_3-R^3 ; with the proviso that when R^7 is H, E' is $-SO_2-$, G is H or alkyl, and when B is present or when B is not present and R^1 is $-C(O)-$, D' may not be C_1-C_{15} alkyl substituted with one substituent selected from $-N(R^3)_2$, $-SR^3$ or $-S(O)_n-R^3$, or substituted with two $-N(R^3)_2$ substituents;

E is selected from Ht; O-Ht; Ht-Ht; Ht fused with Ht; $-O-R^3$; $-N(R^2)(R^3)$; C_1-C_6 alkyl optionally substituted with one or more groups selected from R^4 or Ht; C_2-C_6 alkenyl optionally substituted with one or more groups selected from R^4 or Ht; C_3-C_6 saturated carbocycle optionally substituted with one or more groups selected from R^4 or Ht; or C_5-C_6 unsaturated carbocycle optionally substituted with one or more groups selected from R^4 or Ht;

each R^4 is independently selected from $-OR^2$, $-OR^3$, $-SR^2$, $-SOR^2$, $-SO_2R^2$, $-CO_2R^2$, $-C(O)-NHR^2$, $-C(O)-N(R^2)_2$, $-C(O)-NR^2(OR^2)$, $-S(O)_2-NHR^2$, halo, $-NR^2-C(O)-R^2$, $-N(R^2)_2$ or $-CN$;

each R^7 is independently selected from hydrogen,



wherein each M is independently selected from H, Li, Na, K, Mg, Ca, Ba, $-N(R^2)_4$, C_1-C_{12} -alkyl, C_2-C_{12} -alkenyl, or $-R^6$; wherein 1 to 4 $-CH_2$ radicals of the alkyl or alkenyl group, other than the $-CH_2$ that is bound to Z, is optionally replaced by a heteroatom group selected from O, $S(O)$, $S(O)_2$, or $N(R^2)$; and wherein any hydrogen in said

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alkyl, alkenyl or R^6 is optionally replaced with a substituent selected from oxo, $-OR^2$, $-R^2$, $N(R^2)_2$, $N(R^2)_3$, R^2OH , $-CN$, $-CO_2R^2$, $-C(O)-N(R^2)_2$, $S(O)_2-N(R^2)_2$, $N(R^2)-C(O)-R^2$, $C(O)R^2$, $-S(O)_n-R^2$, OCF_3 , $-S(O)_n-R^6$, $N(R^2)-S(O)_2(R^2)$, halo, $-CF_3$, or $-NO_2$;

M' is H, C_1-C_{12} -alkyl, C_2-C_{12} -alkenyl, or $-R^6$; wherein 1 to 4 $-CH_2$ radicals of the alkyl or alkenyl group is optionally replaced by a heteroatom group selected from O, S, $S(O)$, $S(O)_2$, or $N(R^2)$; and wherein any hydrogen in said alkyl, alkenyl or R^6 is optionally replaced with a substituent selected from oxo, $-OR^2$, $-R^2$, $-N(R^2)_2$, $N(R^2)_3$, $-R^2OH$, $-CN$, $-CO_2R^2$, $-C(O)-N(R^2)_2$, $-S(O)_2-N(R^2)_2$, $-N(R^2)-C(O)-R^2$, $-C(O)R^2$, $-S(O)_n-R^2$, $-OCF_3$, $-S(O)_n-R^6$, $-N(R^2)-S(O)_2(R^2)$, halo, $-CF_3$, or $-NO_2$;

Z is O, S, $N(R^2)_2$, or, when M is not present, H.

Y is P or S;

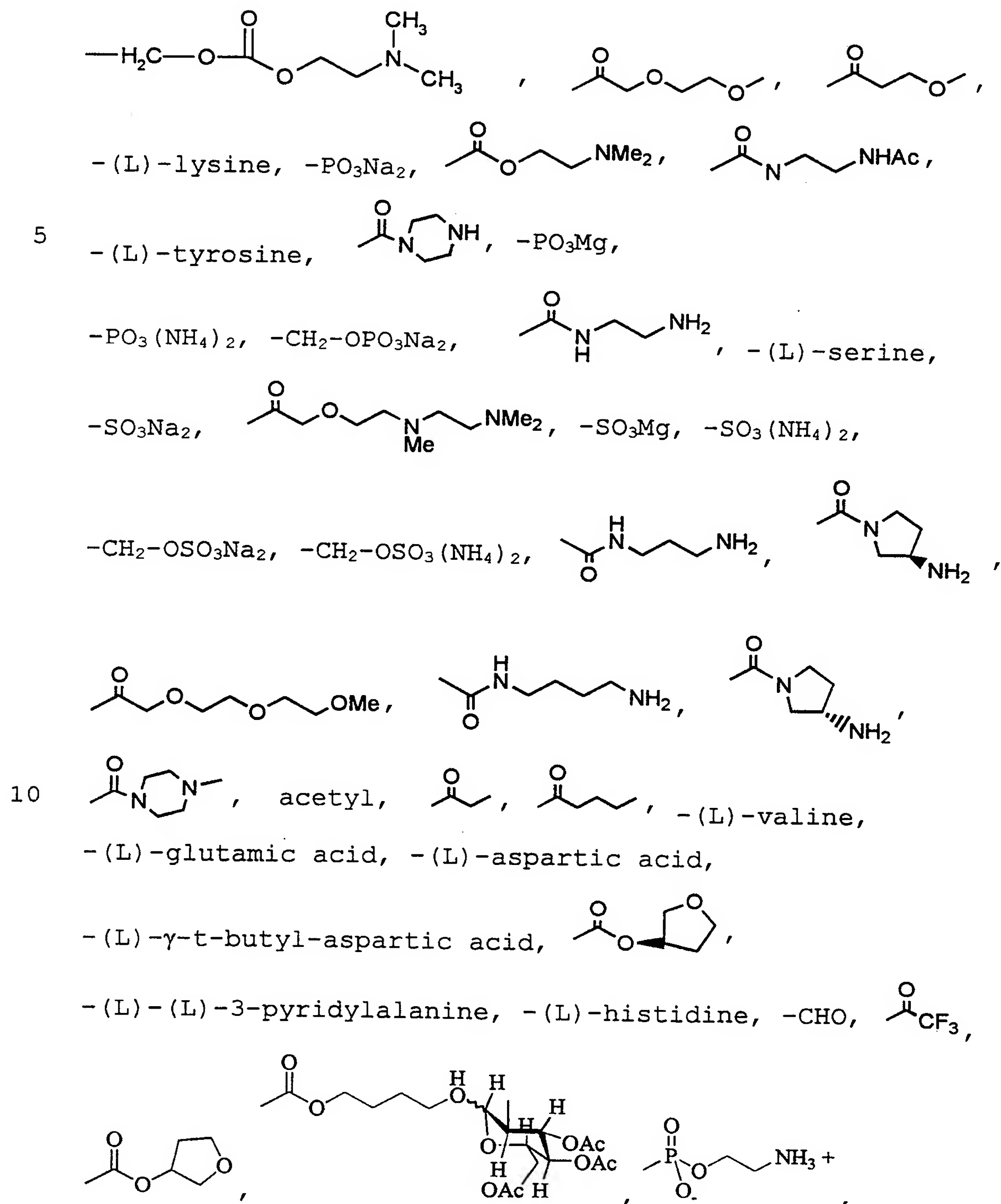
X is O or S;

R^9 is $C(R^2)_2$, O or $N(R^2)$; and wherein when Y is S, Z is not S;

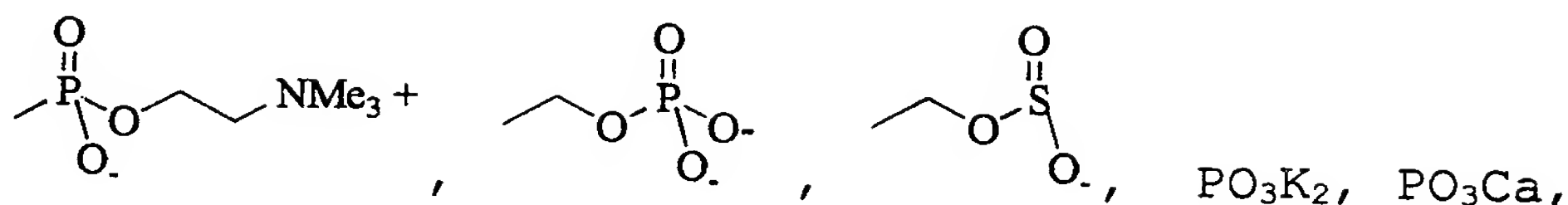
R^6 is a 5-6 membered saturated, partially saturated or unsaturated carbocyclic or heterocyclic ring system, or an 8-10 membered saturated, partially saturated or unsaturated bicyclic ring system; wherein any of said heterocyclic ring systems contains one or more heteroatoms selected from O, N, S, $S(O)_n$ or $N(R^2)$; and wherein any of said ring systems optionally contains 1 to 4 substituents independently selected from OH, C_1-C_4 alkyl, $-O-C_1-C_4$ alkyl or $-O-C(O)-C_1-C_4$ alkyl; and

each R^5 is independently selected from hydrogen, C_1-C_8 alkyl, C_2-C_8 alkenyl, C_2-C_8 alkynyl or Ht, wherein any R^5 , except for hydrogen, is optionally substituted with $-CF_3$, $-PO_3R^3$, azido or halo.

Preferably, at least one R^7 is selected from:



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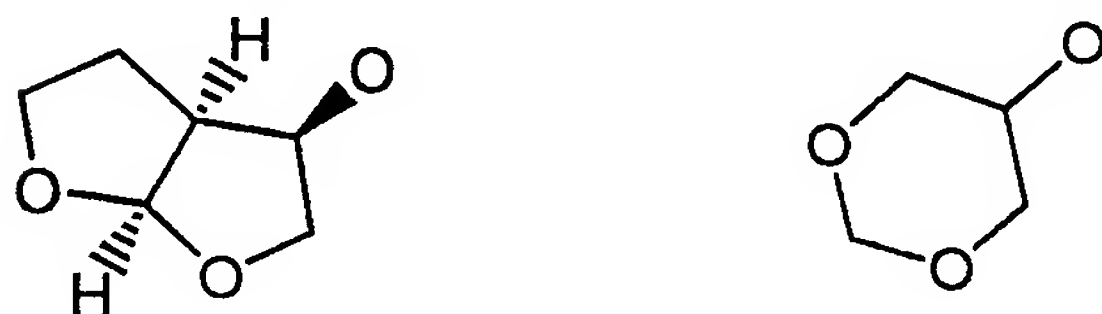
$\text{PO}_3\text{-spermine}$, $\text{PO}_3\text{-(spermidine)}_2$ or $\text{PO}_3\text{-(meglamine)}_2$.

It will be understood by those of skill in the art that component M or M' in the formulae set forth herein will have either a covalent, a covalent/
 5 zwitterionic, or an ionic association with either Z or R⁹ depending upon the actual choice for M or M'. When M or M' is hydrogen, alkyl, alkenyl, or R⁶, M or M' is covalently bound to R⁹ or Z. If M is a mono- or bivalent
 10 metal or other charged species (i.e., NH_4^+), there is an ionic interaction between M and Z and the resulting compound is a salt.

When x is 0 in (M)_x, Z may be a charged species. When that occurs, the other M may be oppositely charged
 15 to produce a 0 net charge on the molecule. Alternatively, the counter ion may be located elsewhere in the molecule.

According to another preferred embodiment, E' is SO_2 .

20 According to yet another preferred embodiment, A-(B)_x is R'-C(O), wherein R' is selected from any of the R' groups indicated in Tables 1 and 2, below. More preferably, R' is selected from:



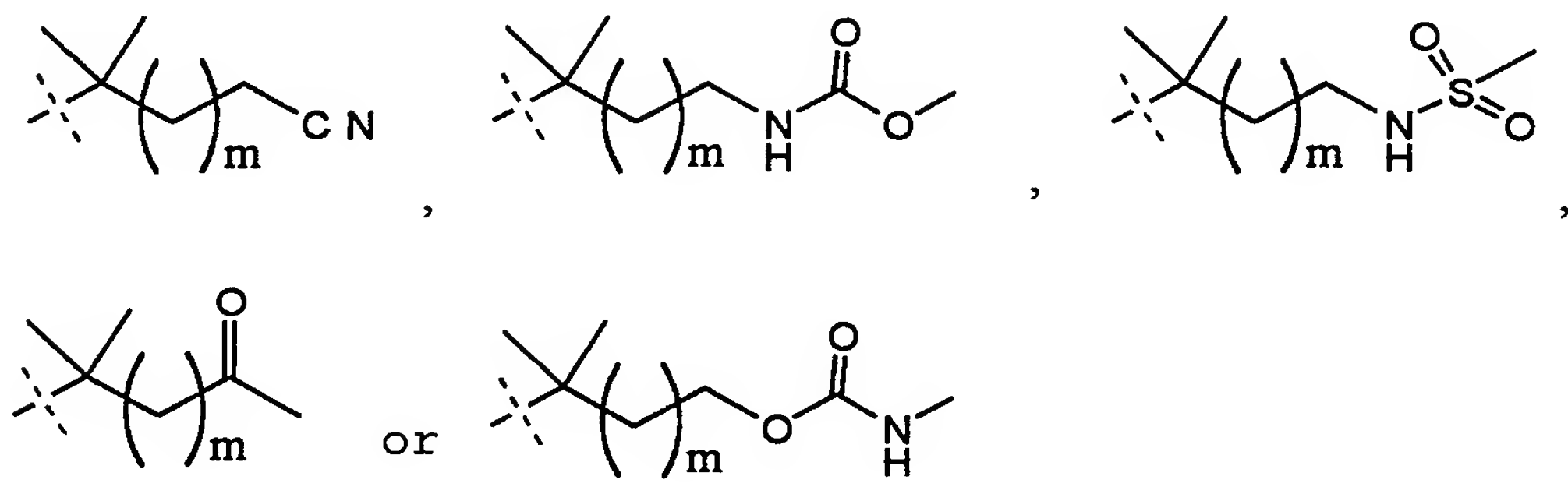
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or

In another preferred embodiment, D' is $-\text{CH}_2\text{-R}''$, wherein R'' is selected from any of the R'' groups

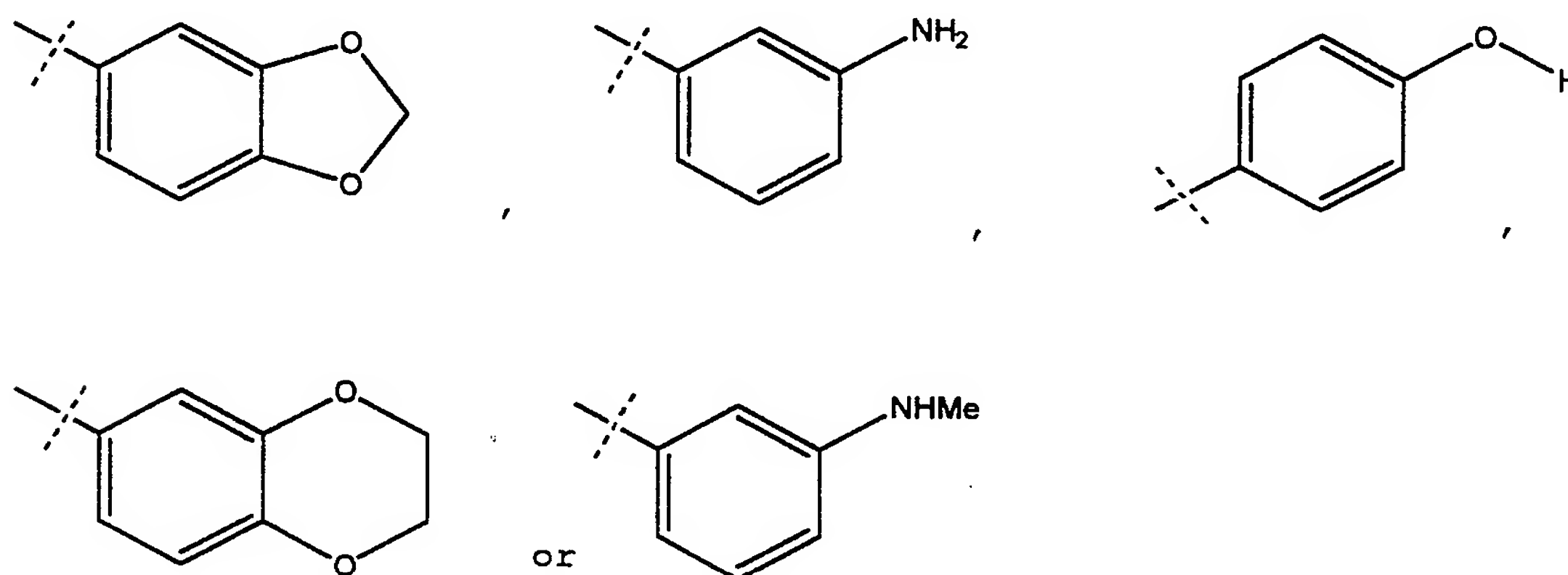
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indicated in Tables 1 and 2, below. More preferably, R'' is selected from:

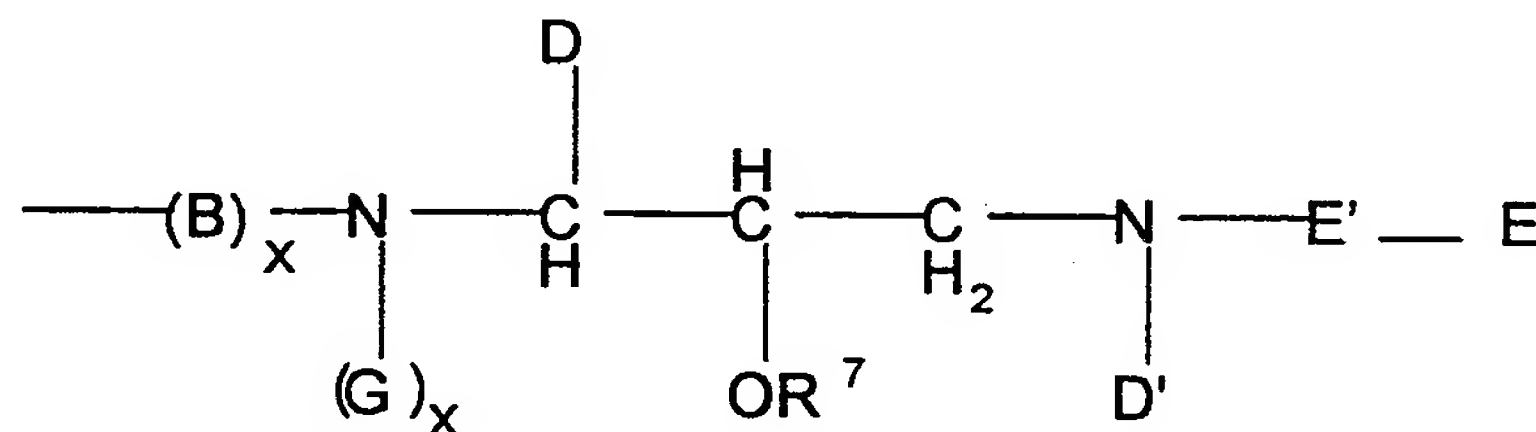


wherein m is 0 to 3.

5 According to another preferred embodiment, E is selected from any of the E groups indicated in Tables 1 and 2, below. More preferably, E is selected from:



10 A more preferred compound of this invention is formula IA:



(IA)

wherein:

E' is -CO- or -SO₂-;

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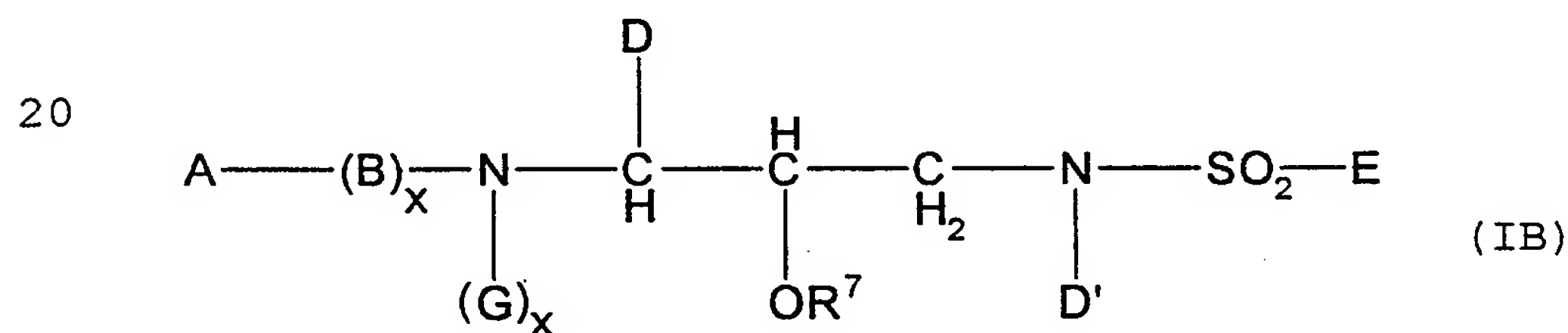
D' is selected from C₁₋₁₅ alkyl, C₂₋₁₅ alkenyl or C₂₋₁₅ alkynyl, wherein D' is substituted with one to two -CN groups and is optionally substituted with C₃₋₈ cycloalkyl; and

5 A, B, D, E, G, R⁷, and x are defined as above for formula I.

In one embodiment of the more preferred compound of this invention represented by formula IA, D' is selected from C₁₋₁₅ alkyl or C₂₋₁₅ alkenyl; each of which is substituted with one to two -CN groups and each of which is optionally substituted with C₃₋₈ cycloalkyl.

In another embodiment of the more preferred compound of this invention represented by formula IA, D' is C₂₋₁₅ alkynyl which is substituted with one to two -CN groups and each of which is optionally substituted with C₃₋₈ cycloalkyl.

Another preferred compound of this invention is represented by formula IB:



wherein:

D' is selected from C₁₋₁₅ alkyl, C₂₋₁₅ alkenyl or C₂₋₁₅ alkynyl, each of which contains one or more substituents selected from oxo, halo, -CF₃, -OCF₃, -NO₂, azido, -SH, -SR³, -N(R³)-N(R³)₂, -O-N(R³)₂, -(R³)N-O-(R³), -N(R³)₂, -CO₂R³, -C(O)-N(R³)₂, -S(O)_n-N(R³)₂, -N(R³)-C(O)-R³, -N(R³)-C(O)-N(R³)₂, -N(R³)-C(O)-S(R³), -C(O)-R³, -S(O)_n-R³, -N(R³)-S(O)_n(R³), -N(R³)-S(O)_n-N(R³)₂, -S-NR³-C(O)R³, -C(S)N(R³)₂, -C(S)R³, -NR³-C(O)OR³, -O-C(O)OR³, -O-

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$C(O)N(R^3)_2$, $-NR^3-C(S)R^3$, $=N-OH$, $=N-OR^3$, $=N-N(R^3)_2$, $=NR^3$,
 $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$, $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-$
 $C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-$
 $C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-OR^3$, $-N(R^3)-C[=N-CN]-OR^3$,
5 $-N(R^3)-C[=N-CN]-(R^3)_2$, $-OC(O)R^3$, $-OC(S)R^3$, $-OC(O)N(R^3)_2$,
 $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-N(R^3)_2$, $O-C(O)N(OR^3)(R^3)$,
 $N(R^3)-N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-$
 $OC(O)R^3$, $-OC(S)N(R^3)_2$, $-OC(S)N(R^3)(R^3)$, or PO_3-R^3 ; with the
proviso that when R^7 is H, E' is $-SO_2$, G is H or alkyl,
10 and when B is present or when B is not present and R^1 is
 $-C(O)-$, D' may not be C_1-C_{15} alkyl substituted with one
substituent selected from $-N(R^3)_2$, $-SR^3$ or $-S(O)_n-R^3$, or
substituted with two $-N(R^3)_2$ substituents; and

15 A, B, D, E, G, R^7 , and x are defined as above for
formula I.

In one embodiment of the more preferred
compound of this invention represented by formula IB, D'
is selected from C_1-C_{15} alkyl or C_2-C_{15} alkenyl, each of
which contains one or more substituents selected from
20 oxo, halo, $-CF_3$, $-OCF_3$, $-NO_2$, azido, $-N(R^3)-N(R^3)_2$, $-O-$
 $N(R^3)_2$, $-(R^3)N-O-(R^3)$, $-N(R^3)_2$, $-N(R^3)-C(O)-N(R^3)_2$, $-N(R^3)-$
 $C(O)-S(R^3)$, $-C(O)-R^3$, $-S(O)_n-R^3$, $-N(R^3)-S(O)_n(R^3)$, $-N(R^3)-$
 $S(O)_n-N(R^3)_2$, $-S-NR^3-C(O)R^3$, $-C(S)N(R^3)_2$, $-C(S)R^3$, $-NR^3-$
 $C(O)OR^3$, $-O-C(O)OR^3$, $-O-C(O)N(R^3)_2$, $-NR^3-C(S)R^3$, $=N-OH$, $=N-$
25 OR^3 , $=N-N(R^3)_2$, $=NR^3$, $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$,
 $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-$
 $C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-$
 OR^3 , $-N(R^3)-C[=N-CN]-OR^3$, $-N(R^3)-C[=N-CN]-(R^3)_2$, $-OC(O)R^3$,
 $-OC(S)R^3$, $-OC(O)N(R^3)_2$, $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-$
30 $N(R^3)_2$, $O-C(O)N(OR^3)(R^3)$, $N(R^3)-N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$,
 $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $-OC(S)N(R^3)_2$, $-OC(S)N(R^3)(R^3)$,
or PO_3-R^3 ; C_2-C_{15} alkynyl which contains one or more

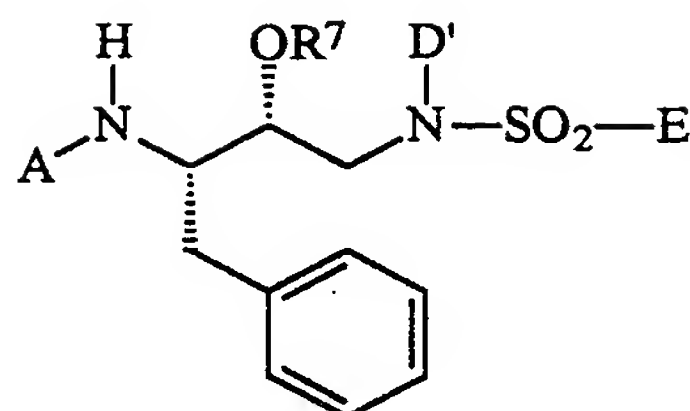
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substituents selected from oxo, halo, $-\text{CF}_3$, $-\text{OCF}_3$, $-\text{NO}_2$, azido, $-\text{SH}$, $-\text{SR}^3$, $-\text{N}(\text{R}^3)-\text{N}(\text{R}^3)_2$, $-\text{O}-\text{N}(\text{R}^3)_2$, $-(\text{R}^3)\text{N}-\text{O}-(\text{R}^3)$, $-\text{N}(\text{R}^3)_2$, $-\text{CO}_2\text{R}^3$, $-\text{C}(\text{O})-\text{N}(\text{R}^3)_2$, $-\text{S}(\text{O})_n-\text{N}(\text{R}^3)_2$, $-\text{N}(\text{R}^3)-\text{C}(\text{O})-\text{R}^3$, $-\text{N}(\text{R}^3)-\text{C}(\text{O})-\text{N}(\text{R}^3)_2$, $-\text{N}(\text{R}^3)-\text{C}(\text{O})-\text{S}(\text{R}^3)$, $-\text{C}(\text{O})-\text{R}^3$, $-\text{S}(\text{O})_n-\text{R}^3$, $-\text{N}(\text{R}^3)-\text{S}(\text{O})_n(\text{R}^3)$, $-\text{N}(\text{R}^3)-\text{S}(\text{O})_n-\text{N}(\text{R}^3)_2$, $-\text{S}-\text{NR}^3-\text{C}(\text{O})\text{R}^3$, $-\text{C}(\text{S})\text{N}(\text{R}^3)_2$, $-\text{C}(\text{S})\text{R}^3$, $-\text{NR}^3-\text{C}(\text{O})\text{OR}^3$, $-\text{O}-\text{C}(\text{O})\text{OR}^3$, $-\text{O}-\text{C}(\text{O})\text{N}(\text{R}^3)_2$, $-\text{NR}^3-\text{C}(\text{S})\text{R}^3$, $=\text{N}-\text{OH}$, $=\text{N}-\text{OR}^3$, $=\text{N}-\text{N}(\text{R}^3)_2$, $=\text{NR}^3$, $=\text{NNR}^3\text{C}(\text{O})\text{N}(\text{R}^3)_2$, $=\text{NNR}^3\text{C}(\text{O})\text{OR}^3$, $=\text{NNR}^3\text{S}(\text{O})_n-\text{N}(\text{R}^3)_2$, $-\text{NR}^3-\text{C}(\text{S})\text{OR}^3$, $-\text{NR}^3-\text{C}(\text{S})\text{N}(\text{R}^3)_2$, $-\text{NR}^3-\text{C}[=\text{N}(\text{R}^3)]-\text{N}(\text{R}^3)_2$, $-\text{N}(\text{R}^3)-\text{C}[=\text{N}-\text{NO}_2]-\text{N}(\text{R}^3)_2$, $-\text{N}(\text{R}^3)-\text{C}[=\text{N}-\text{NO}_2]-\text{OR}^3$, $-\text{N}(\text{R}^3)-\text{C}[=\text{N}-\text{CN}]-\text{OR}^3$, $-\text{N}(\text{R}^3)-\text{C}[=\text{N}-\text{CN}]-\text{N}(\text{R}^3)_2$, $-\text{OC}(\text{O})\text{R}^3$, $-\text{OC}(\text{S})\text{R}^3$, $-\text{OC}(\text{O})\text{N}(\text{R}^3)_2$, $-\text{C}(\text{O})\text{N}(\text{R}^3)-\text{N}(\text{R}^3)_2$, $-\text{O}-\text{C}(\text{O})\text{N}(\text{R}^3)-\text{N}(\text{R}^3)_2$, $\text{O}-\text{C}(\text{O})\text{N}(\text{OR}^3)(\text{R}^3)$, $\text{N}(\text{R}^3)-\text{N}(\text{R}^3)\text{C}(\text{O})\text{R}^3$, $\text{N}(\text{R}^3)-\text{OC}(\text{O})\text{R}^3$, $\text{N}(\text{R}^3)-\text{OC}(\text{O})\text{R}^3$, $\text{N}(\text{R}^3)-\text{OC}(\text{O})\text{R}^3$, $-\text{OC}(\text{S})\text{N}(\text{R}^3)_2$, $-\text{OC}(\text{S})\text{N}(\text{R}^3)(\text{R}^3)$, or PO_3-R^3 ; with the proviso that when R^7 is H, E' is $-\text{SO}_2-$, G is H or alkyl, and when B is present or when B is not present and R^1 is $-\text{C}(\text{O})-$, D' may not be C_1-C_{15} alkyl substituted with one substituent selected from $-\text{N}(\text{R}^3)_2$ or $-\text{S}(\text{O})_n-\text{R}^3$, or substituted with two $-\text{N}(\text{R}^3)_2$ substituents.

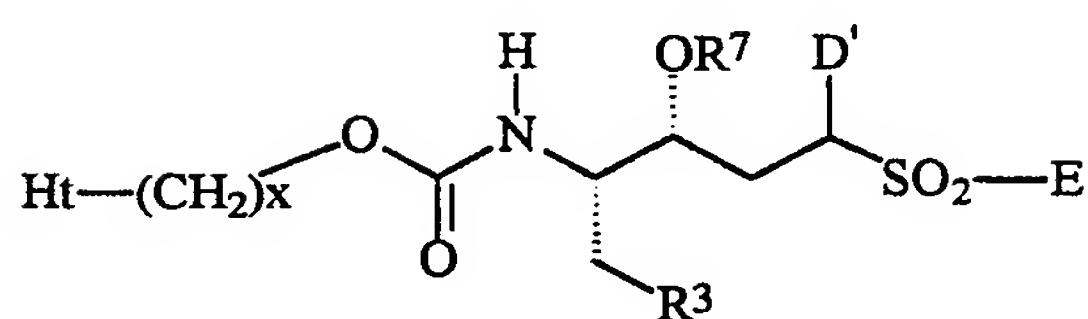
In another embodiment of the more preferred compound of this invention represented by formula IB, D' is selected from C_1-C_{15} alkyl or C_2-C_{15} alkenyl, each of which contains one or more substituents selected from $-\text{SH}$, $-\text{SR}^3$, $-\text{CO}_2\text{R}^3$, $-\text{C}(\text{O})-\text{N}(\text{R}^3)_2$, $-\text{S}(\text{O})_n-\text{N}(\text{R}^3)_2$ or $-\text{N}(\text{R}^3)-\text{C}(\text{O})-\text{R}^3$; with the proviso that when R^7 is H, E' is $-\text{SO}_2-$, G is H or alkyl, and when B is present or when B is not present and R^1 is $-\text{C}(\text{O})-$, D' may not be C_1-C_{15} alkyl substituted with one substituent selected from $-\text{SR}^3$.

More preferred compounds of formula I are those represented by formula II, formula III or formula IV:

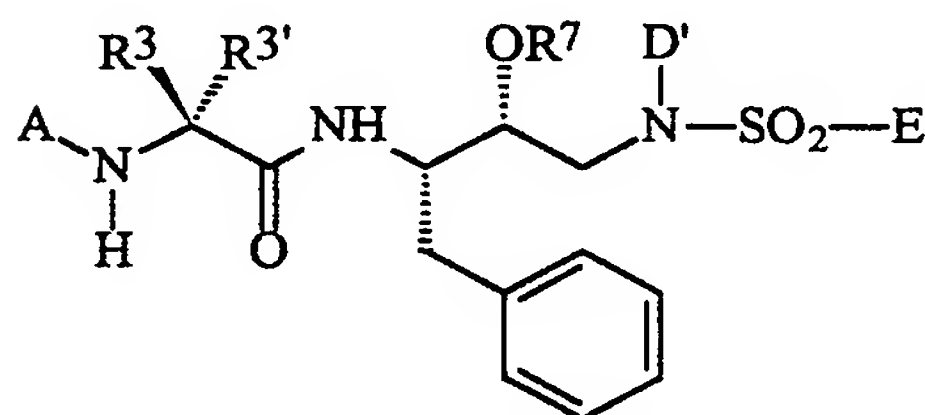
-24-



(II)



(III)



(IV);

- 5 wherein A, D', E, R³, R⁷, Ht and x are as defined for compounds of Formula I, above. For ease of reference, the two R³ moieties present in formula IV have been labeled R³ and R^{3'} and wherein R^{3'} is selected from H, Ht, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₃-C₆ cycloalkyl or C₅-C₆ cycloalkenyl; wherein any member of said R³, except H, is optionally substituted with one or more substituents selected from -OR², -C(O)-NH-R², -S(O)_n-N(R²)(R²), -N(R²)₂, -N(R²)-C(O)-O(R²), -N(R²)-C(O)-N(R²), -N(R²)-C(O)-(R²), -N(R²-OR²)₂, -C(O)-Ht, Ht, -CN, -SR², -CO₂R², or NR²-C(O)-R².

15 For compounds of formula II, more preferred compounds are those wherein:

A is -C(O)Ht;

E is C₆-C₁₀ aryl optionally substituted with one or more substituents selected from oxo, -OR², SR², -R²,
 20 -N(R²)₂, -R²-OH, -CN, -CO₂R², -C(O)-N(R²)₂, -S(O)₂-N(R²)₂,
 -N(R²)-C(O)-R², -C(O)-R², -S(O)_n-R², -OCF₃, -S(O)_n-Q,

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methylenedioxy, $-N(R^2)-S(O)_2(R^2)$, halo, $-CF_3$, $-NO_2$, Q, $-OQ$,
5 $-OR^7$, $-SR^7$, $-R^7$, $-N(R^2)(R^7)$ or $-N(R^7)_2$; or a 5-membered
heterocyclic ring containing one S and optionally
containing N as an additional heteroatom, wherein said
heterocyclic ring is optionally substituted with one to
two groups independently selected from $-CH_3$, R^4 , or Ht.

Another preferred embodiment of compounds of
formula II are those wherein:

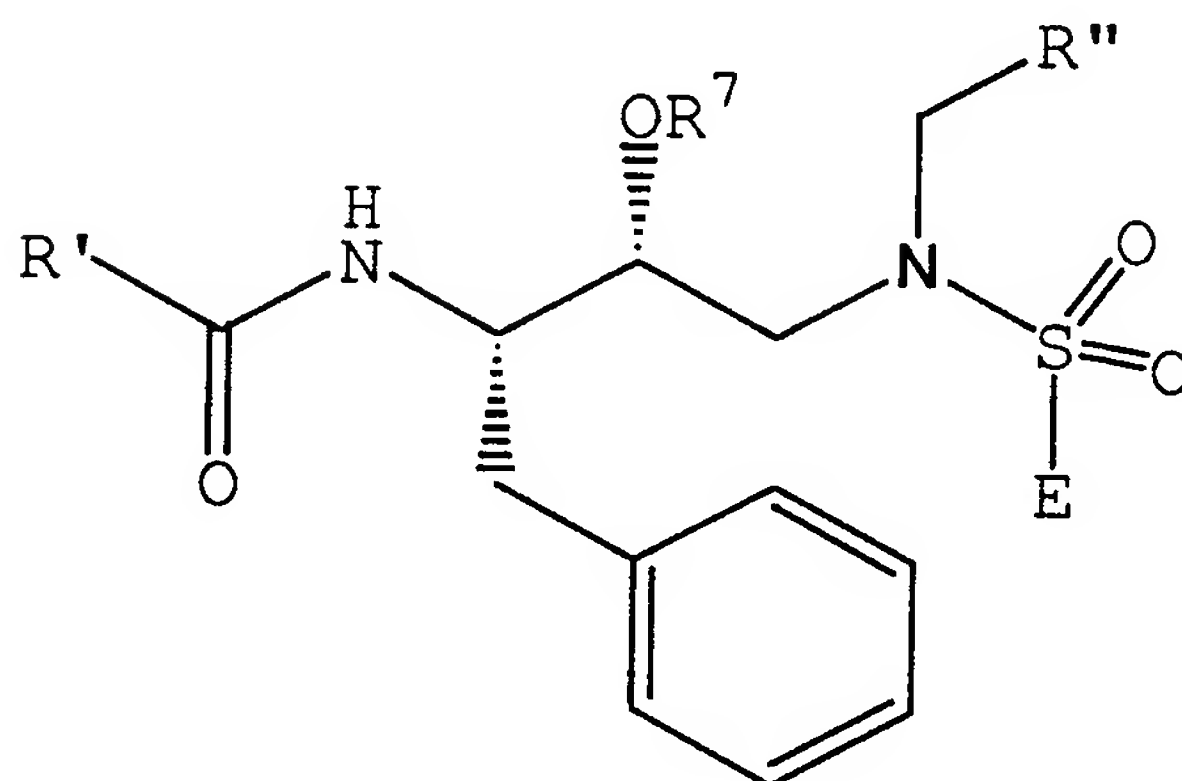
10 E is a 5-membered heterocyclic ring containing one S
and optionally containing N as an additional heteroatom,
wherein said heterocyclic ring is optionally substituted
with one to two groups independently selected from $-CH_3$,
 R^4 , or Ht.

More preferred are compounds of formula II set
15 forth above, wherein R^7 in $-OR^7$ is $-PO(OM)_2$ or
 $C(O)CH_2OCH_2CH_2OCH_2CH_2OCH_3$ and both R^7 in $-N(R^7)_2$ are H,
wherein M is H, Li, Na, K or C_1 - C_4 alkyl; or wherein R^7 in
 $-OR^7$ is $C(O)CH_2OCH_2CH_2OCH_3$, one R^7 in $-N(R^7)_2$ is
 $C(O)CH_2OCH_2CH_2OCH_3$ and the other is H.

20 The compounds according to the invention
contain one or more asymmetric carbon atoms and thus
occur as racemates and racemic mixtures, single
enantiomers, diastereomeric mixtures and individual
diastereomers. All such isomeric forms of these
25 compounds are expressly included in the present
invention. Each stereogenic carbon may be of the R or S
configuration. Although the specific compounds
exemplified in this application may be depicted in a
particular stereochemical configuration, compounds having
30 either the opposite stereochemistry at any given chiral
center or mixtures thereof are also envisioned.

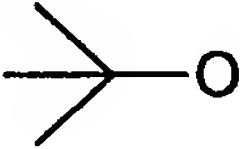
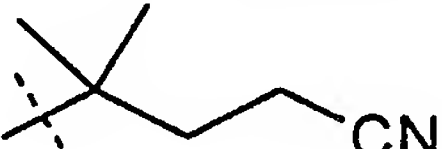
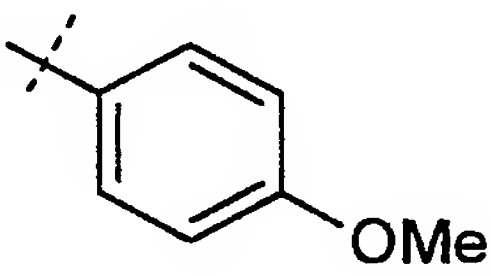
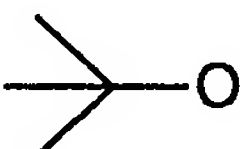
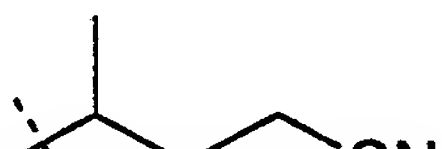
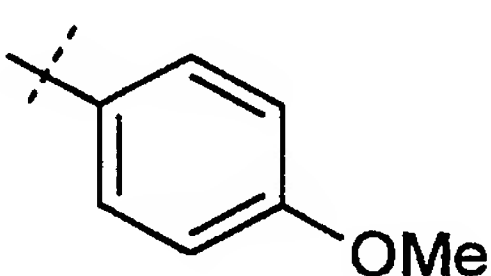

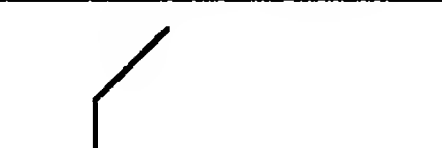
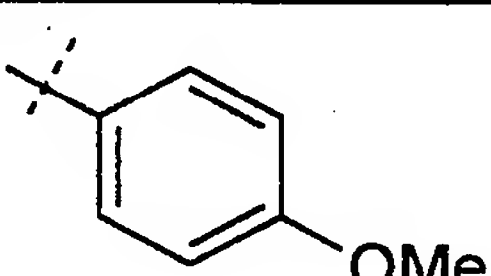
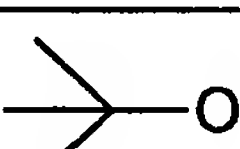
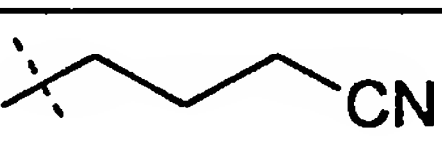
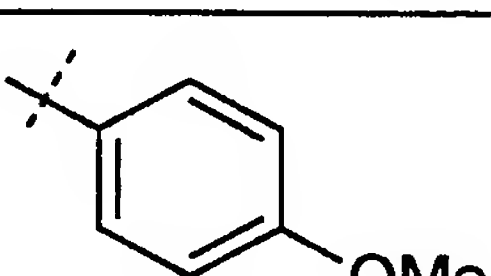

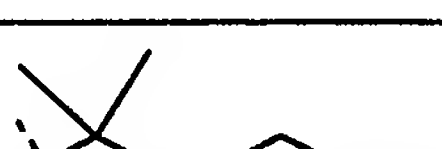
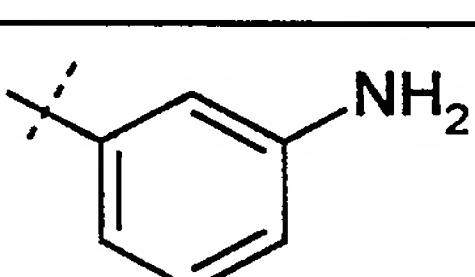
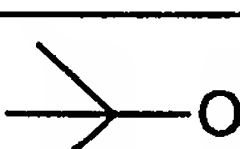

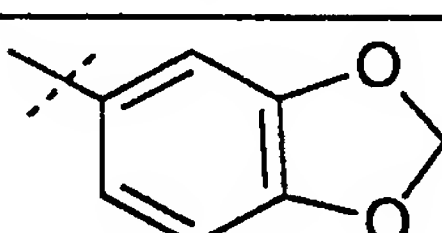
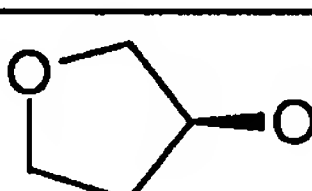

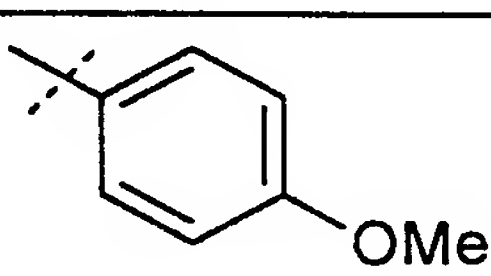
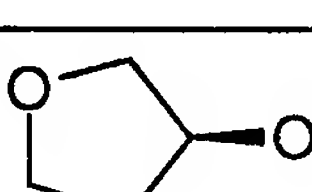
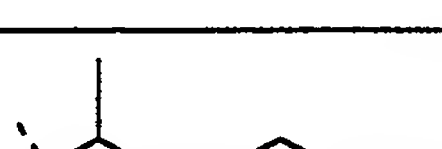
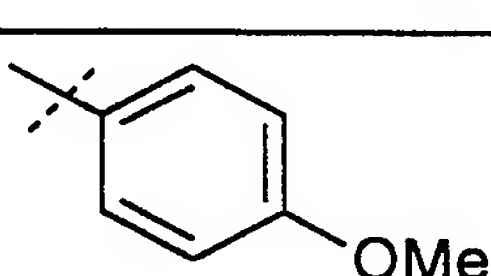
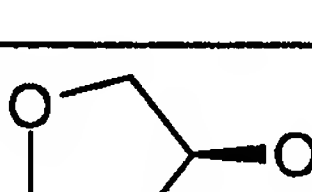
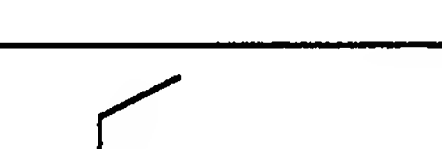
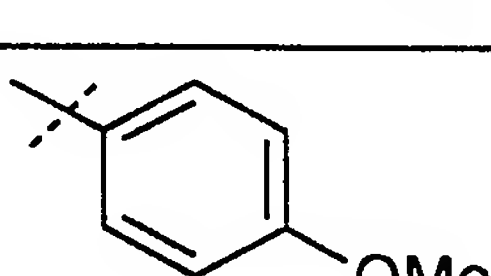
Specific preferred compounds of the present invention are set forth below in Table 1.

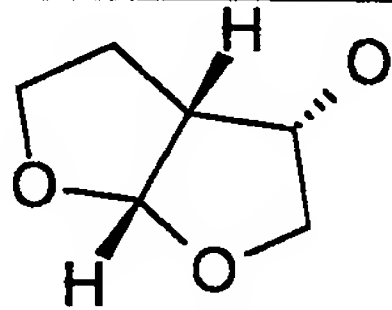
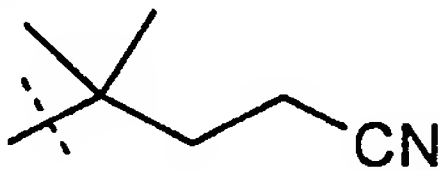
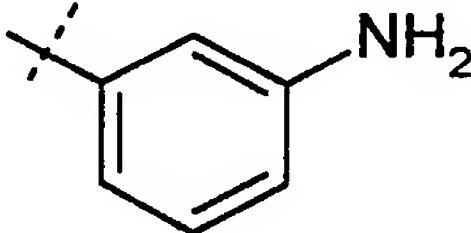
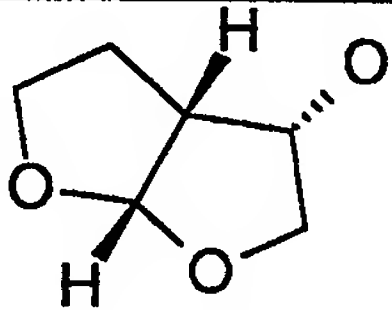
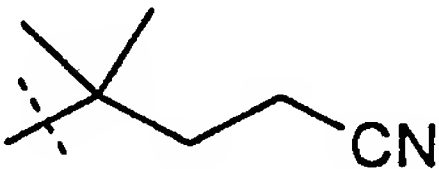
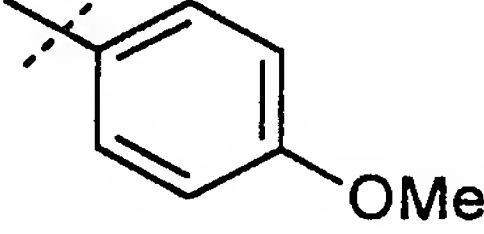
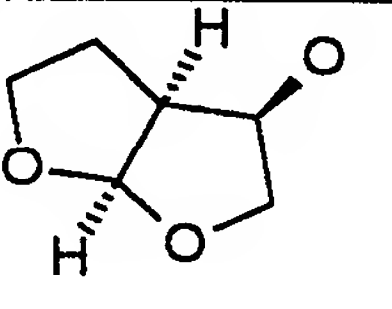
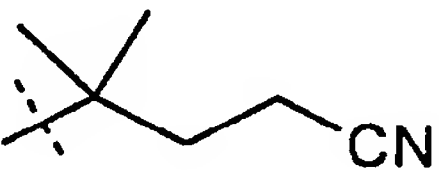
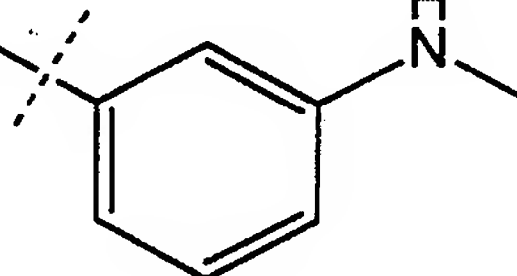
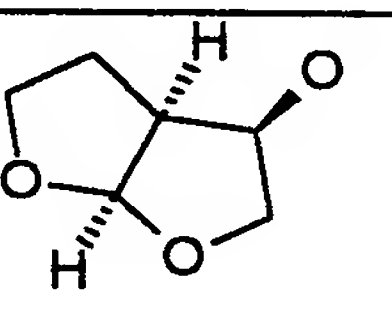
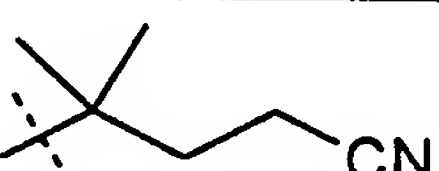
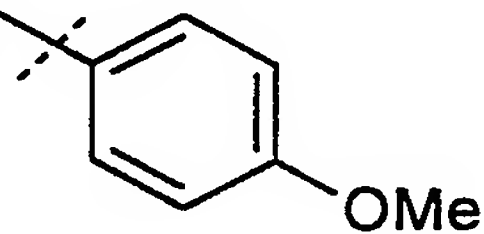
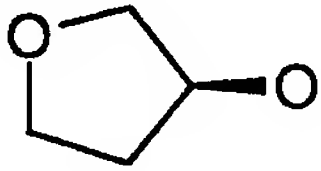
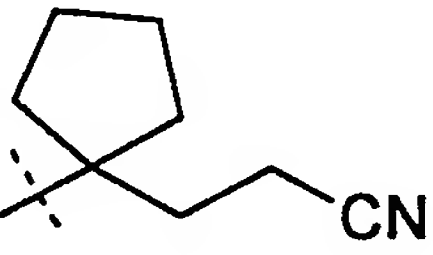
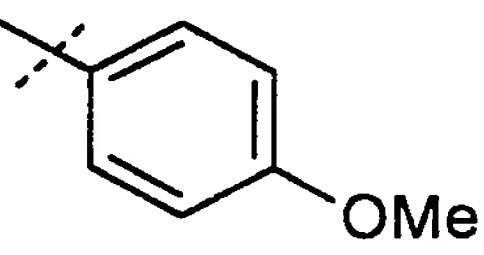
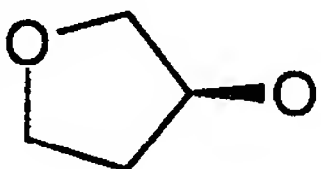
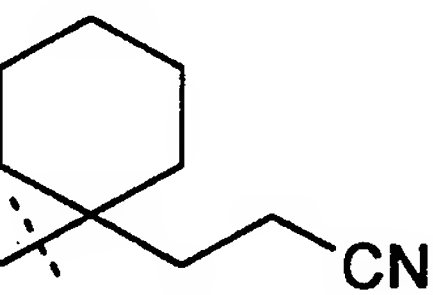
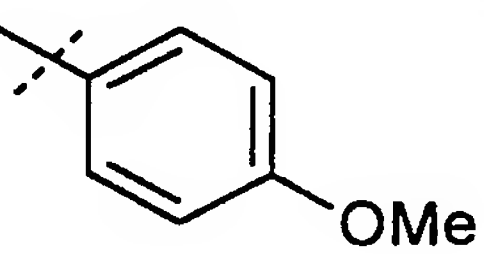
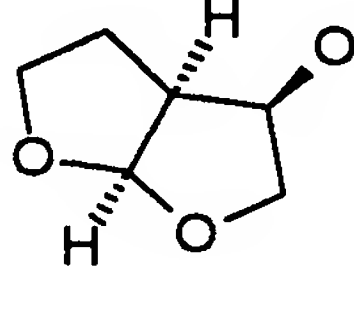
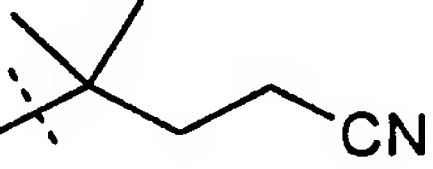
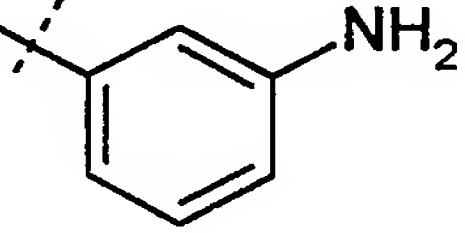
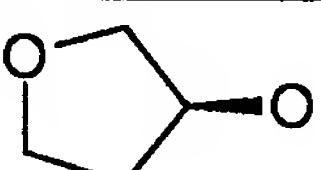
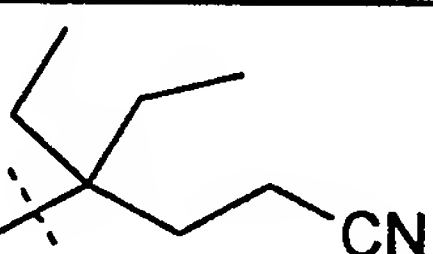
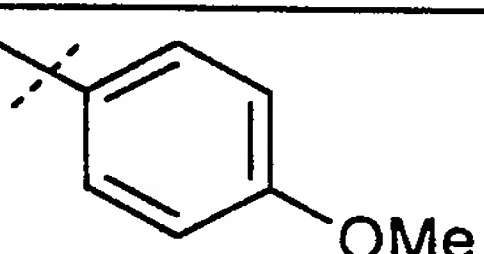
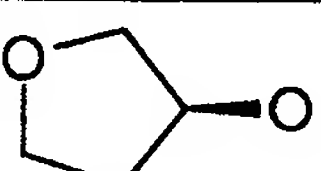

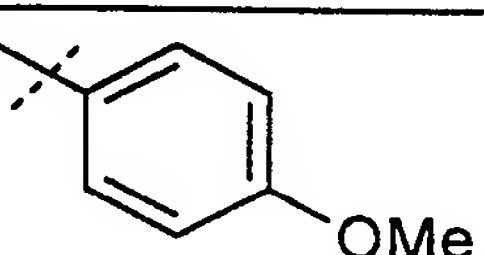
Table 1

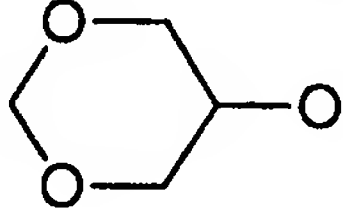
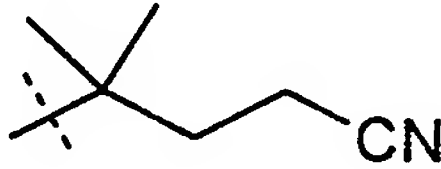
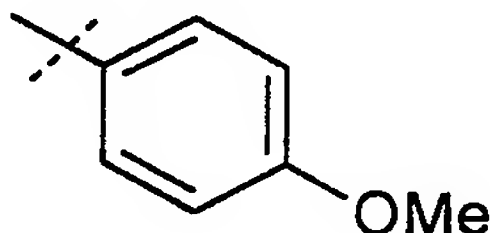
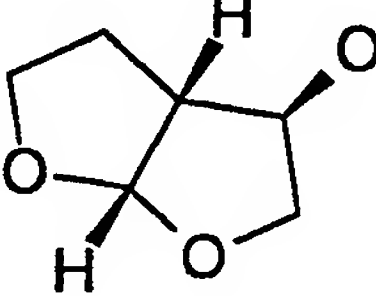
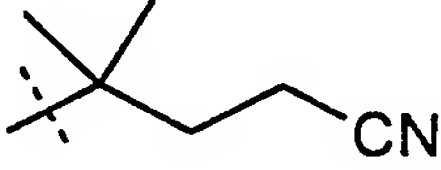
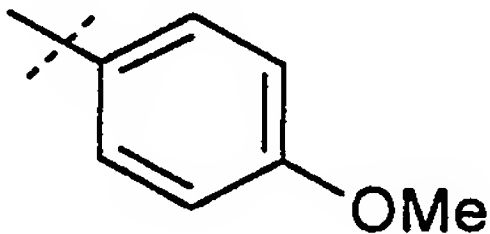
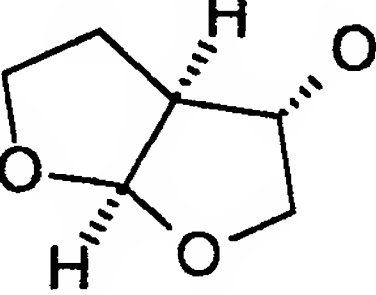
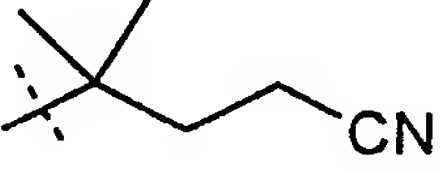
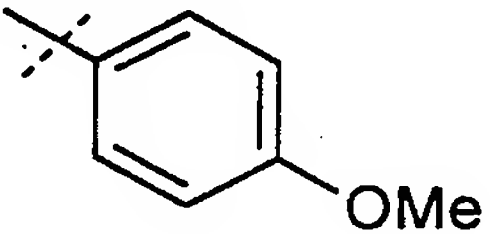
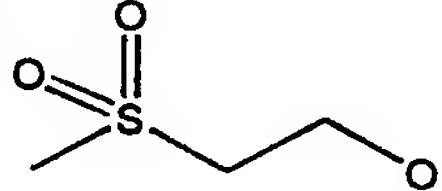
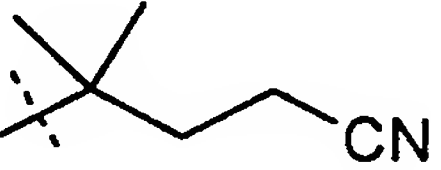
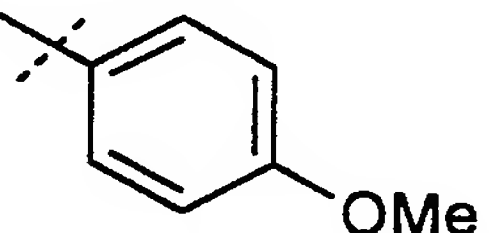
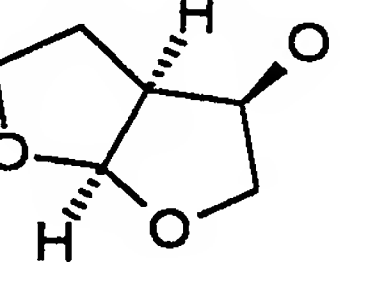
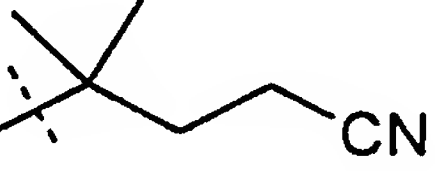
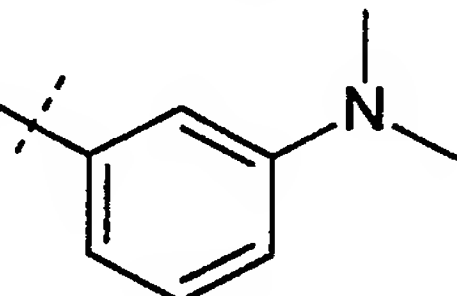
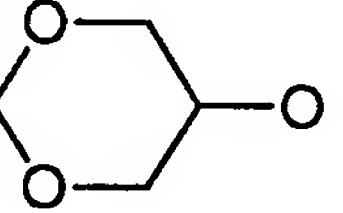
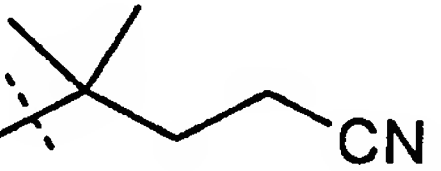
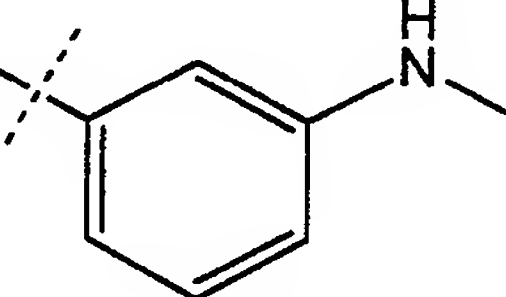
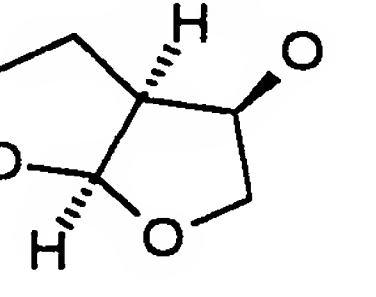
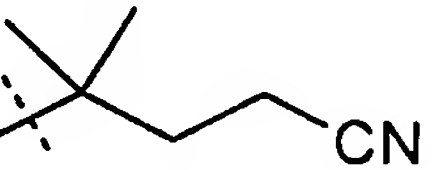
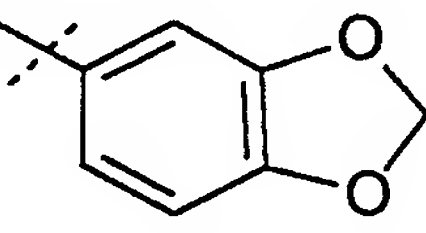
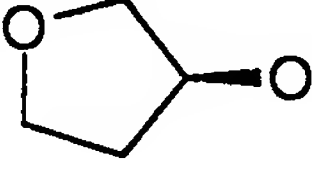
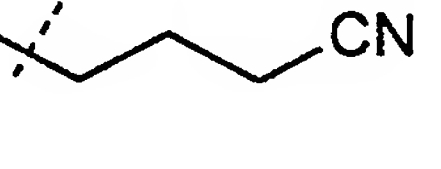
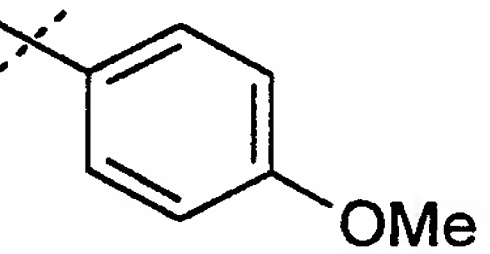


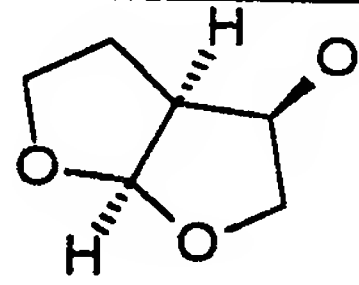
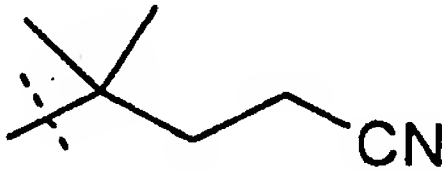
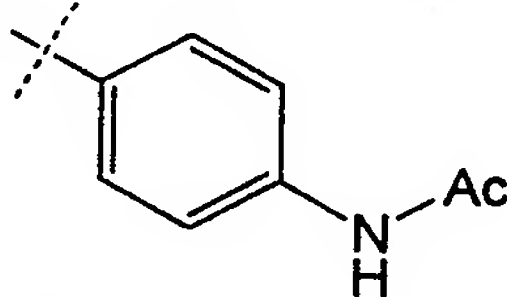
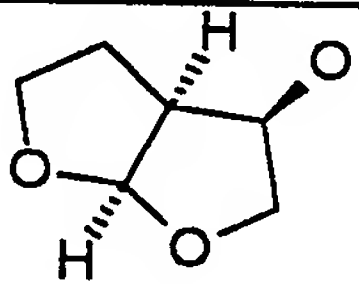
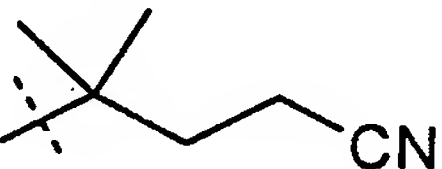
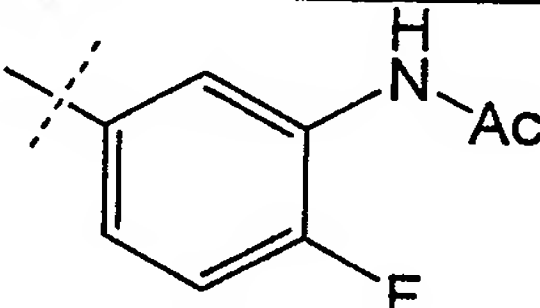
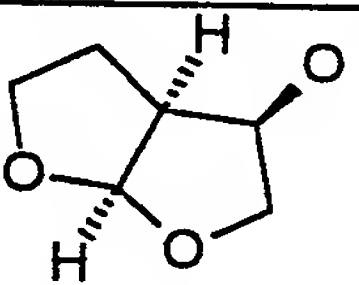
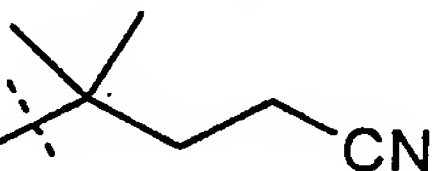
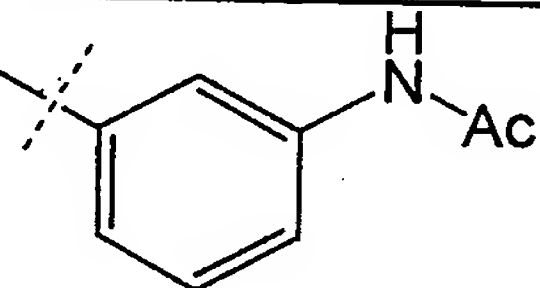
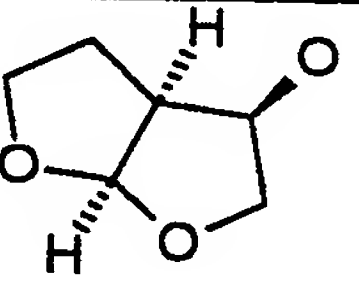
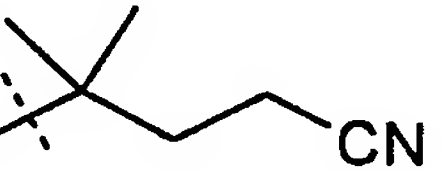
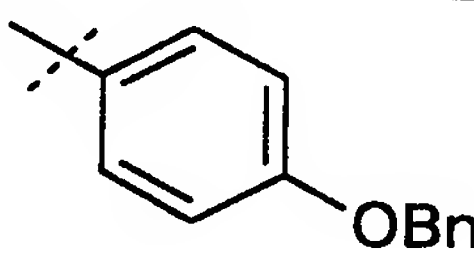
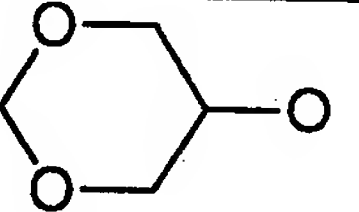
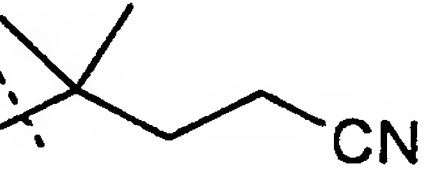
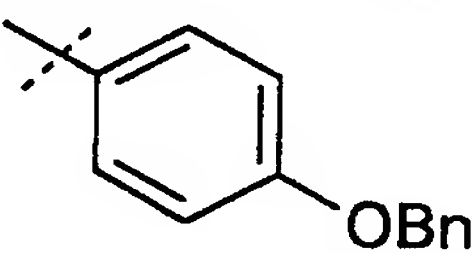
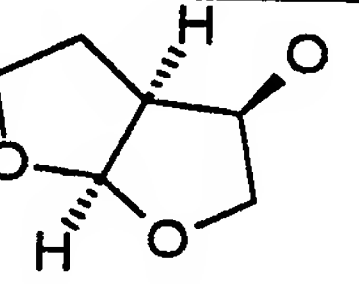
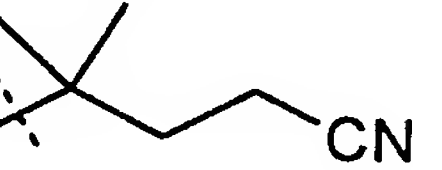
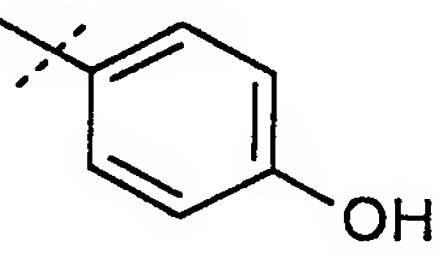
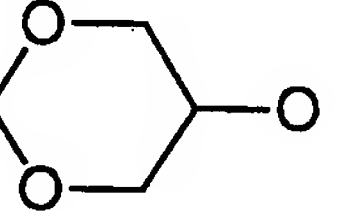
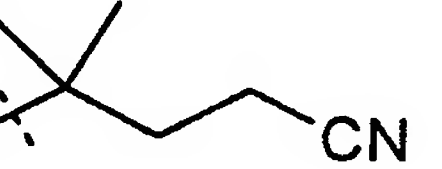
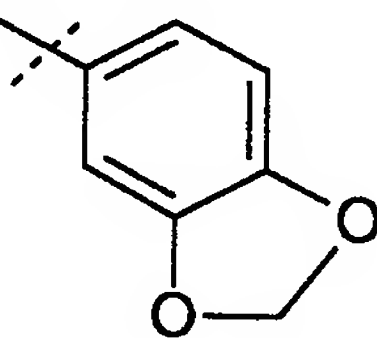
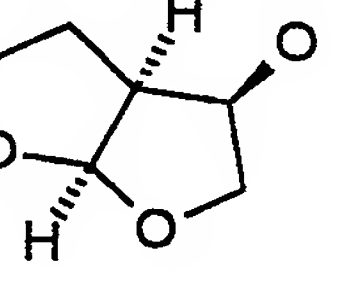
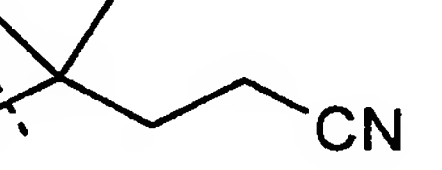
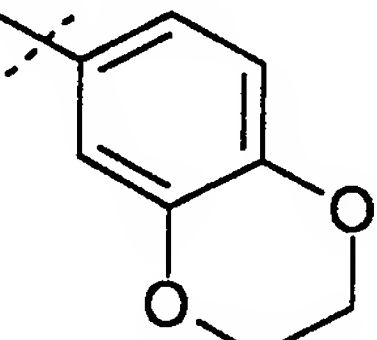
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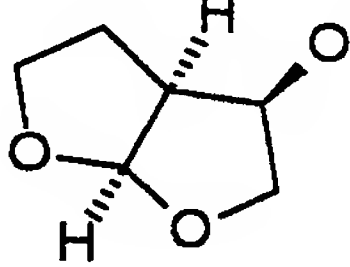
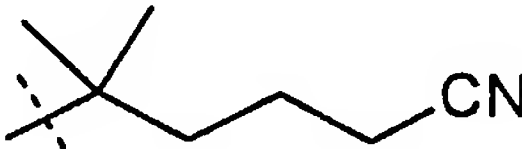
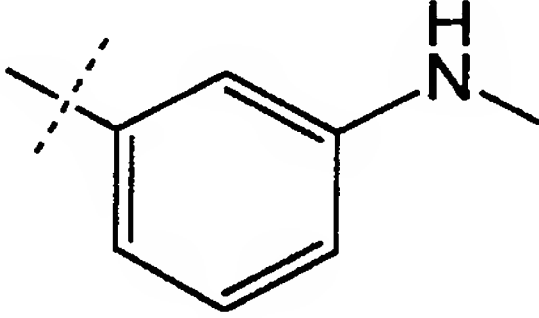
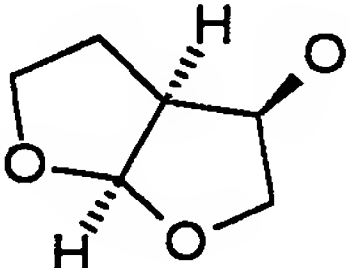
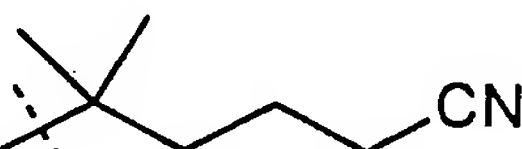
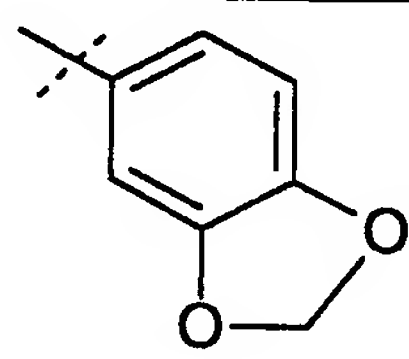
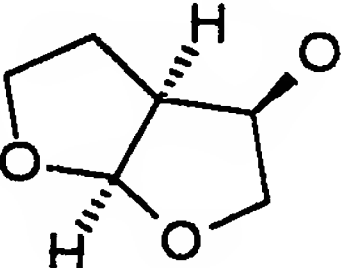
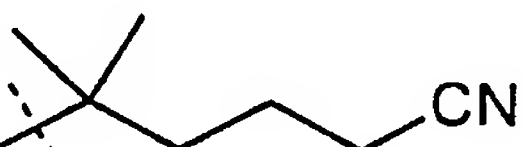
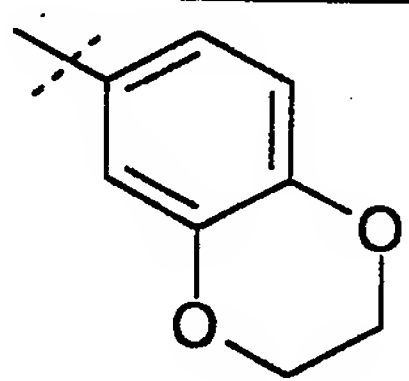
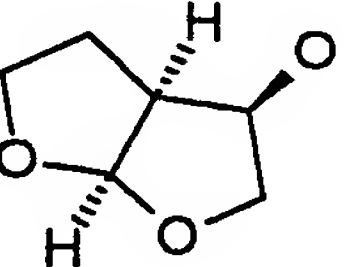
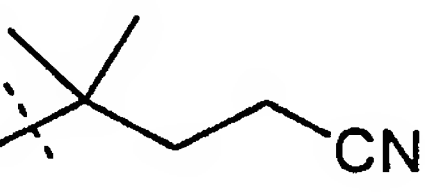
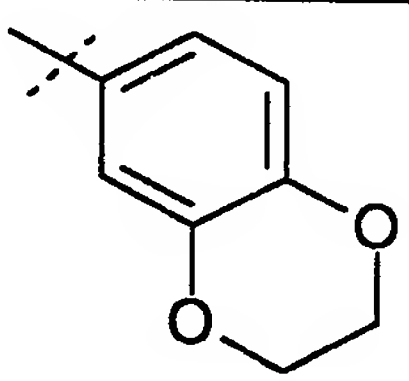
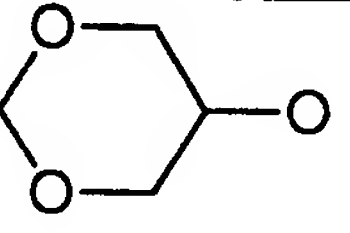
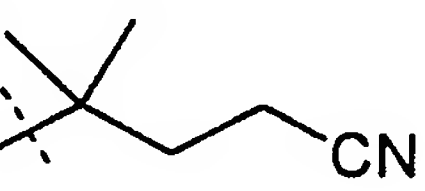
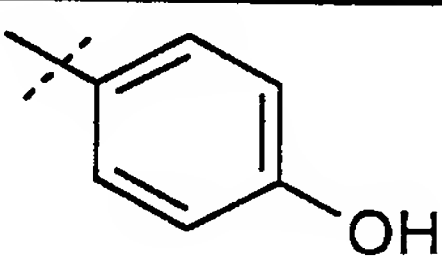
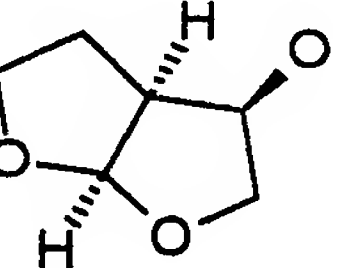
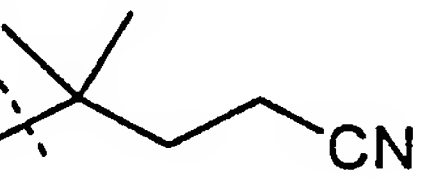
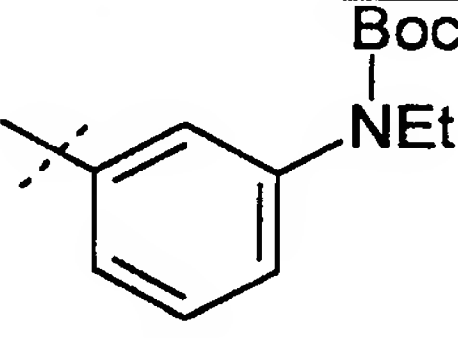
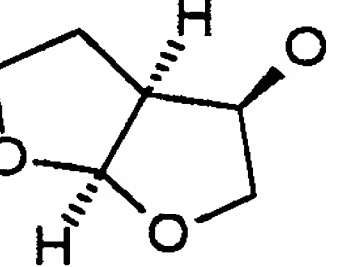
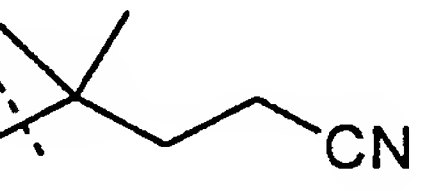
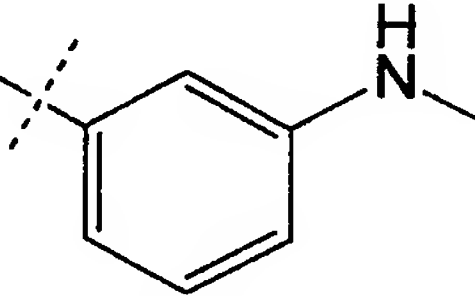
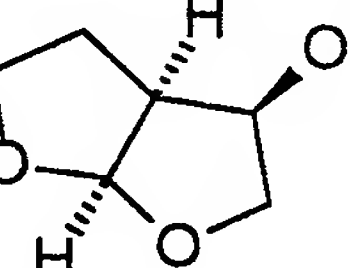
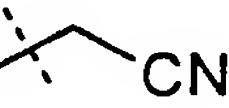
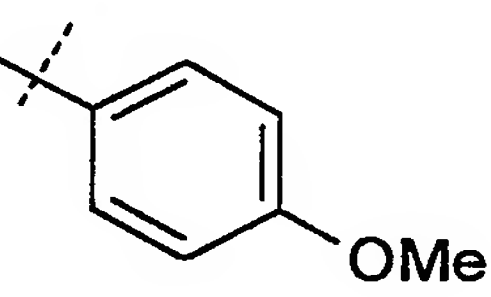
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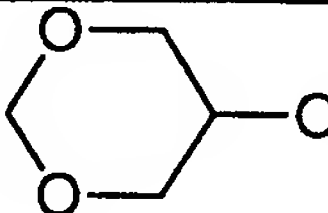
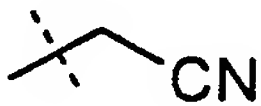
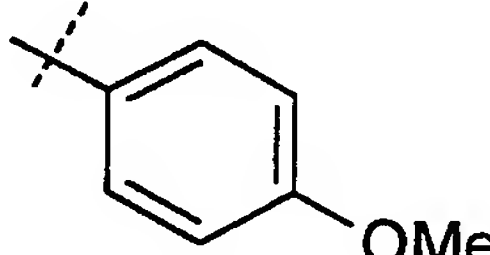
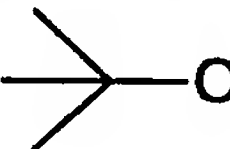
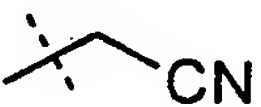
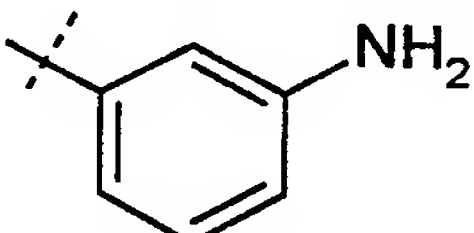
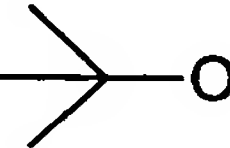
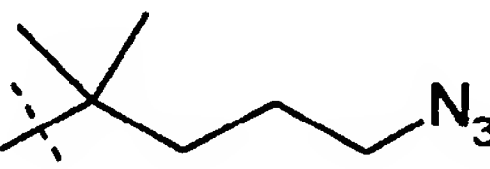
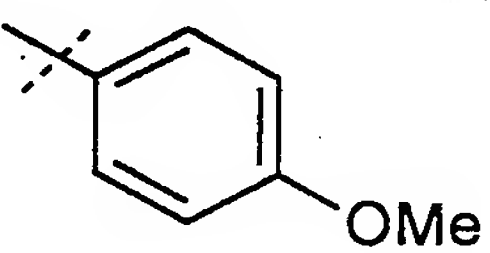
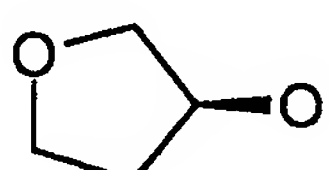
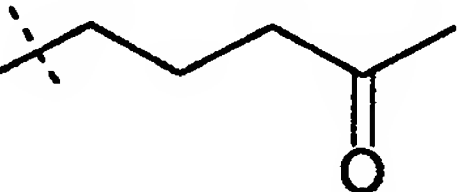
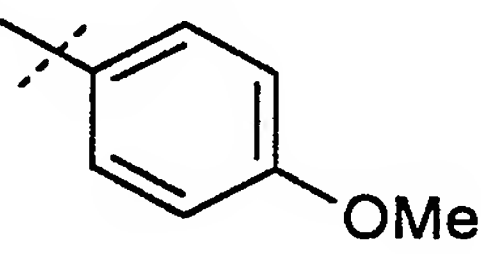
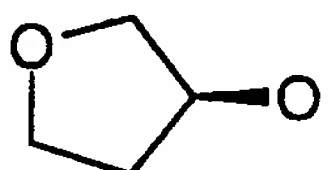
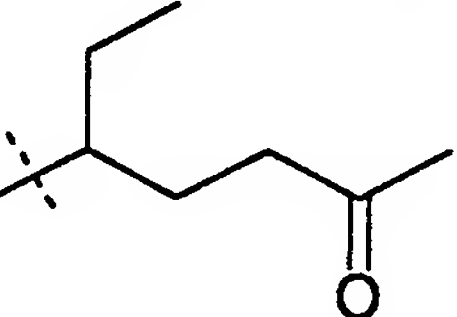
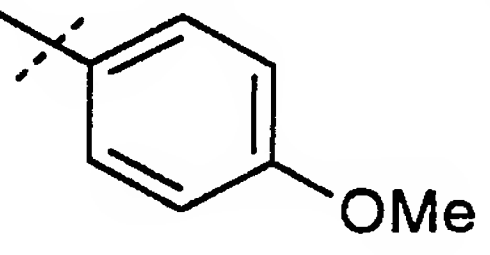

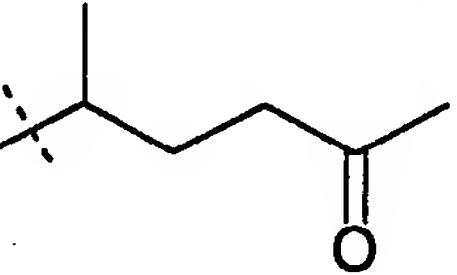
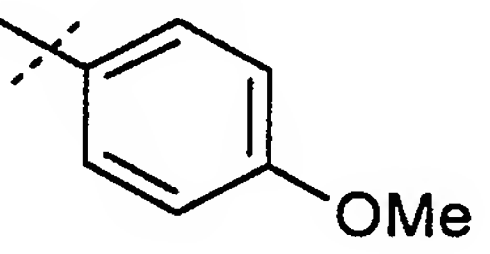
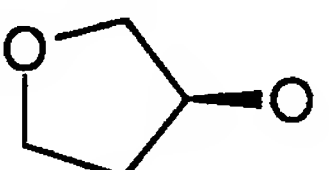
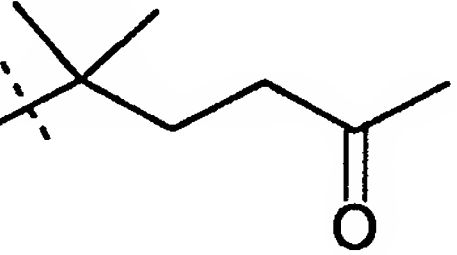
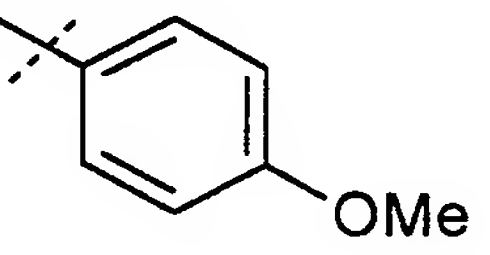
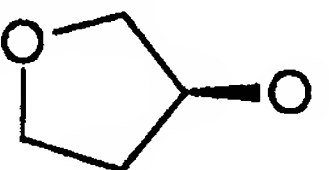
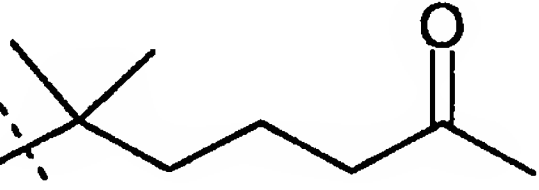
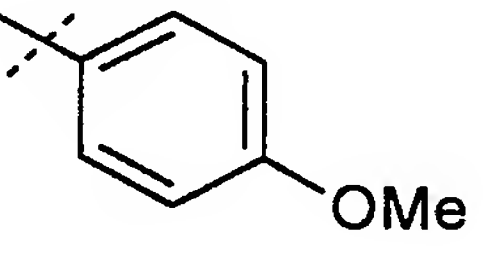
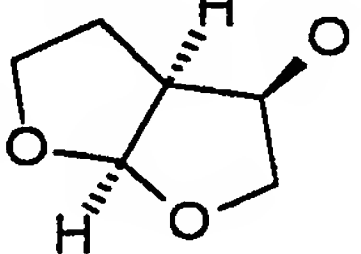
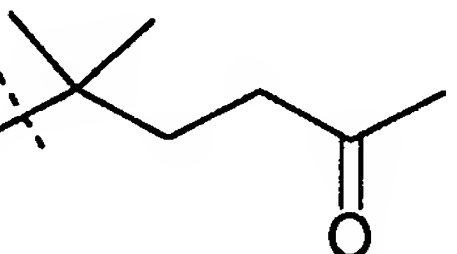
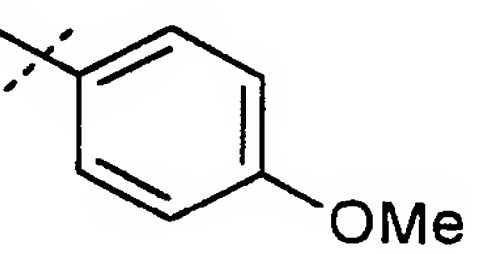
Compound	R'	R''	E
1			
2			
3			
4			
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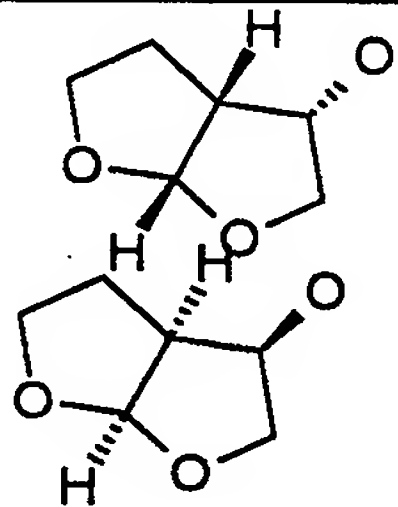
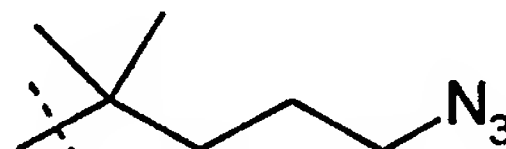
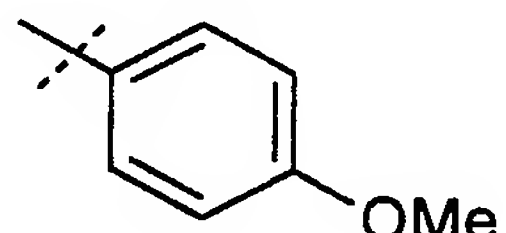
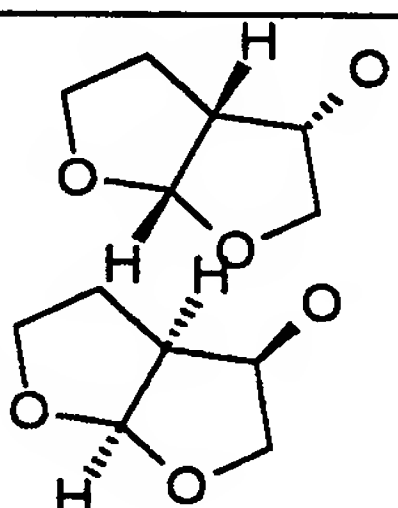
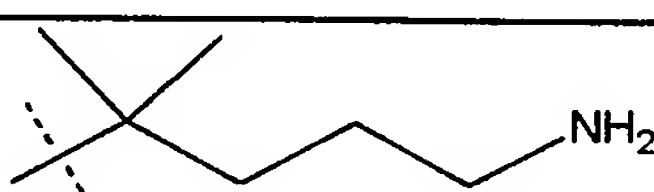
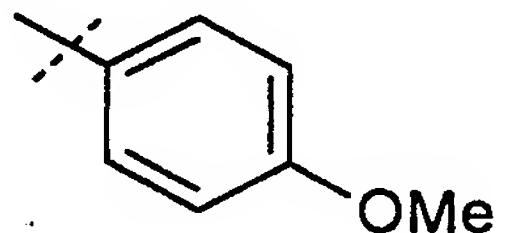
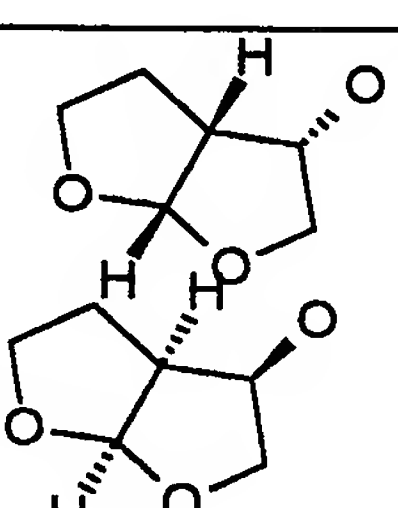
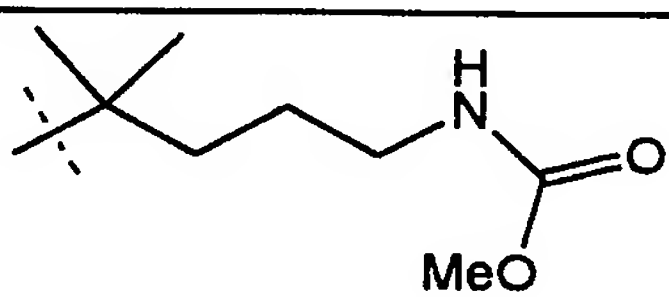
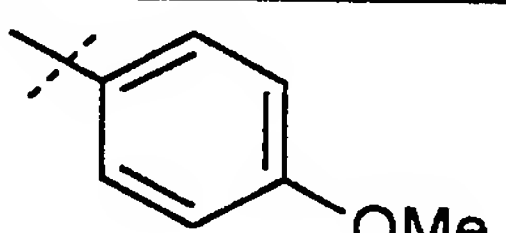
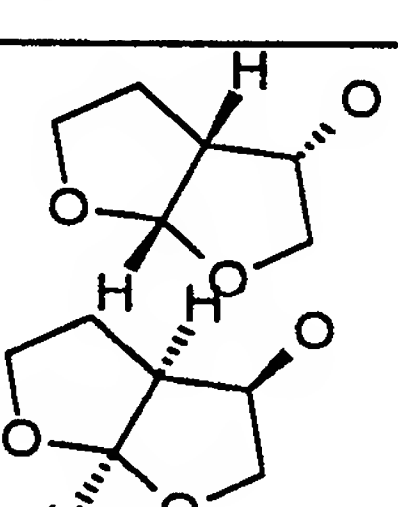
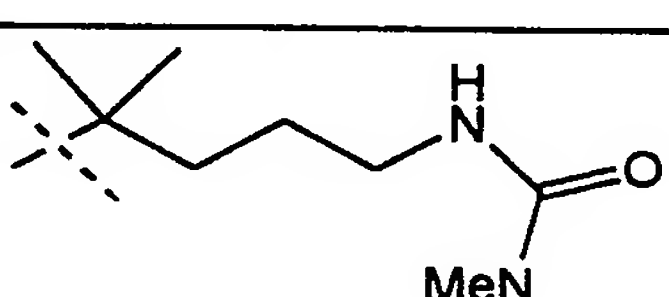
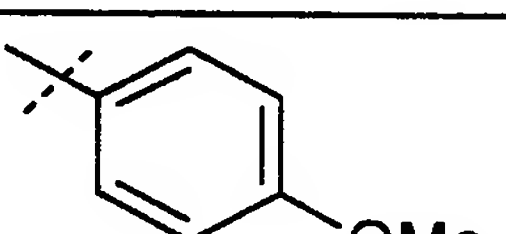
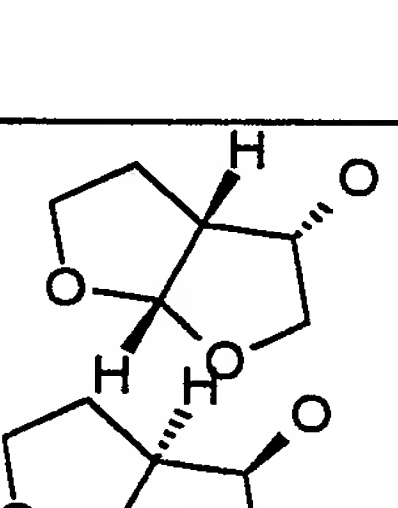
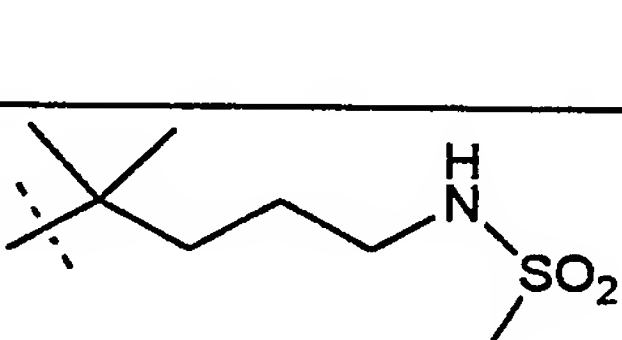

Compound	R'	R''	E
10			
11			
12			
13			
14			
15			
16			
17			
18			

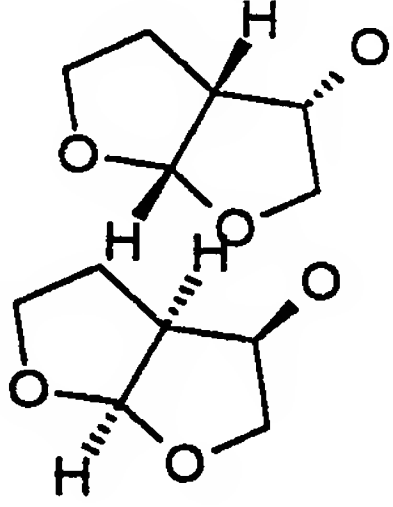
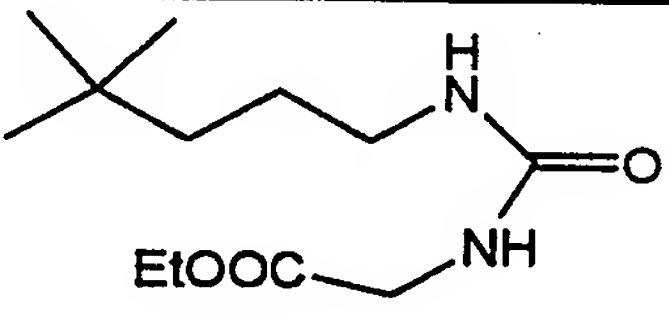
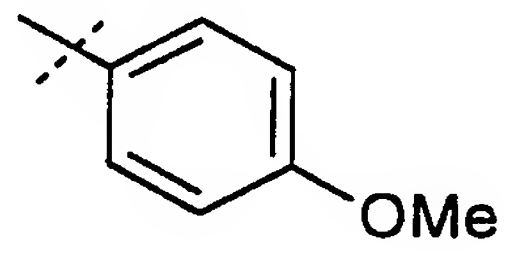
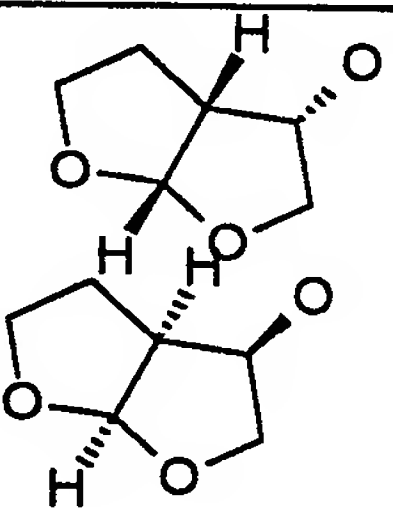
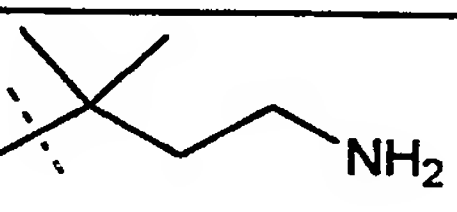
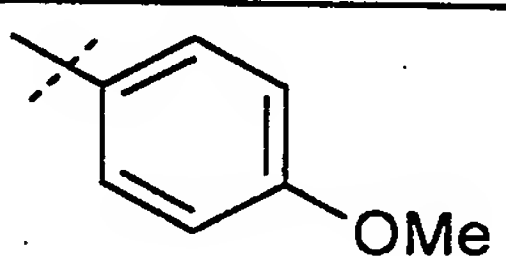
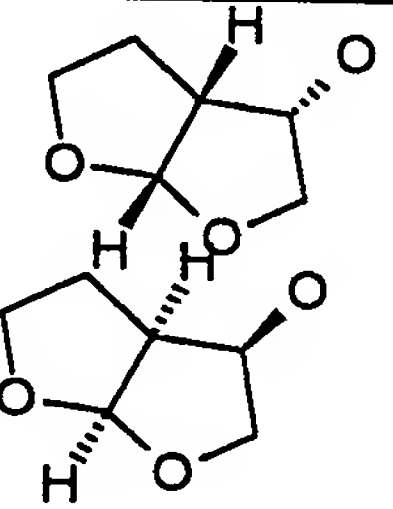
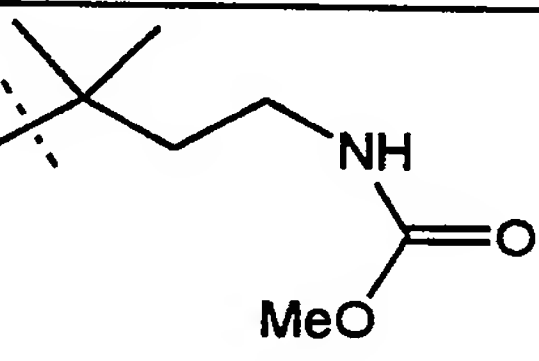
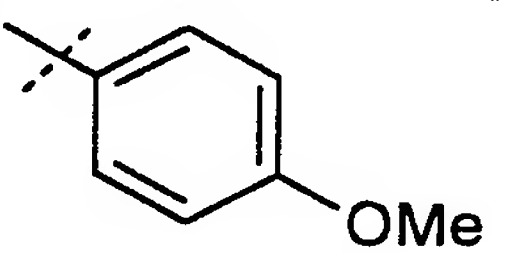
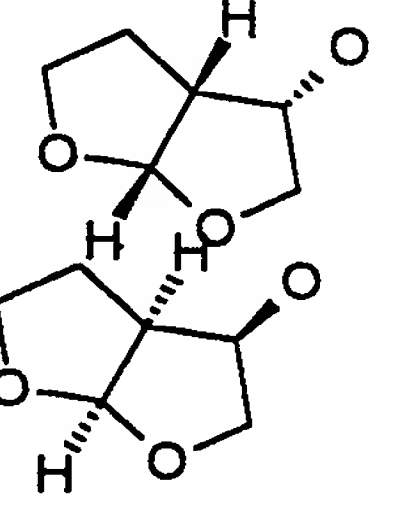
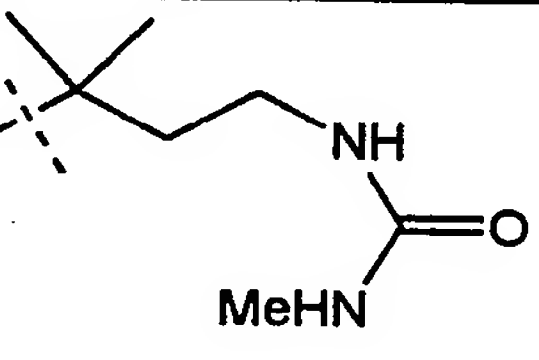
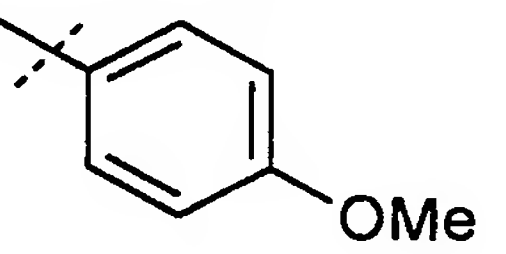
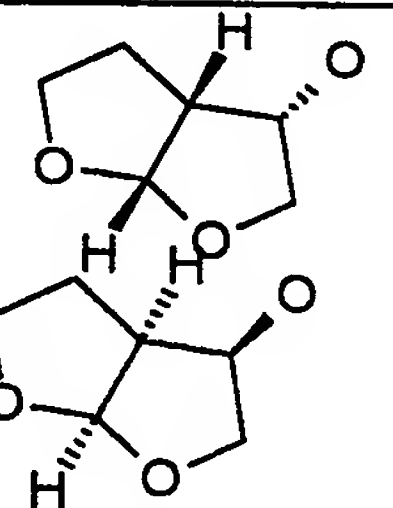
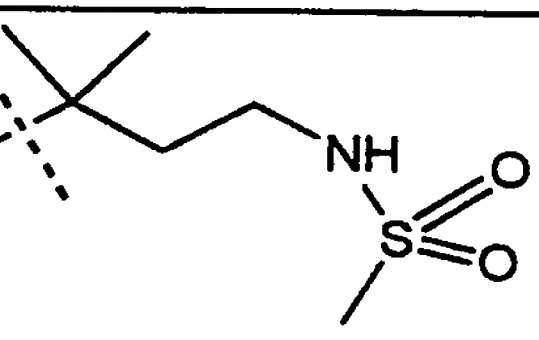
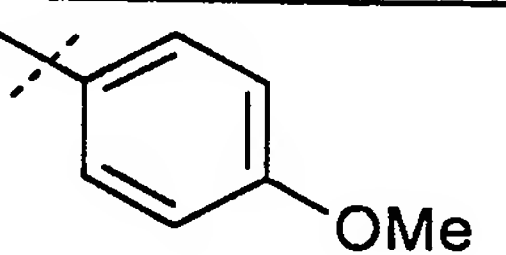
Compound	R'	R''	E
19			
20			
21			
22			
23			
24			
25			
26			

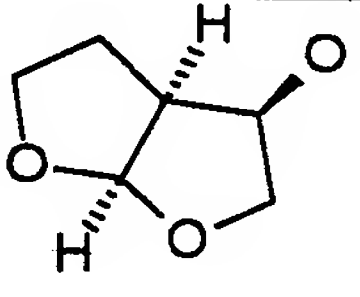
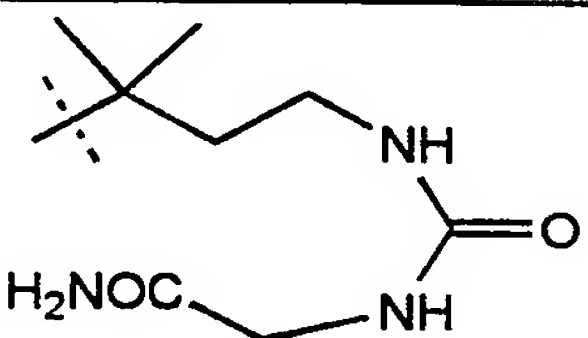
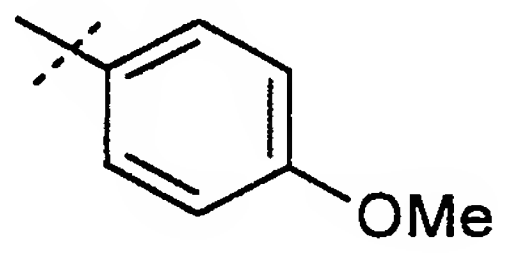
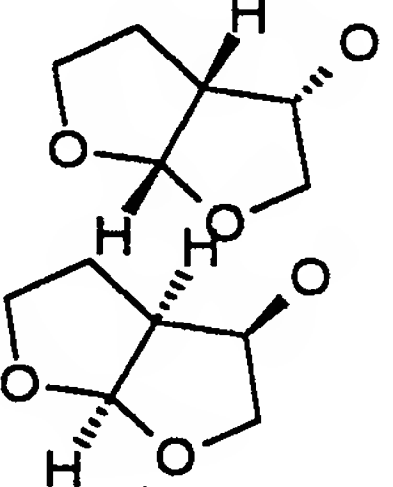
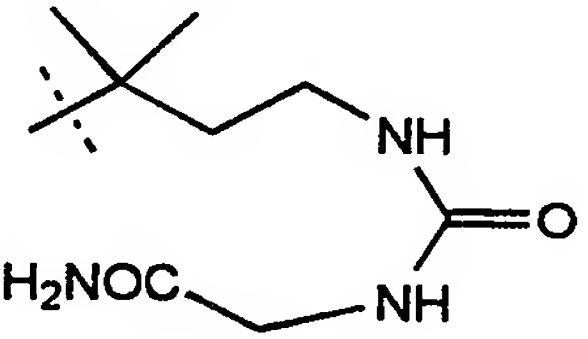
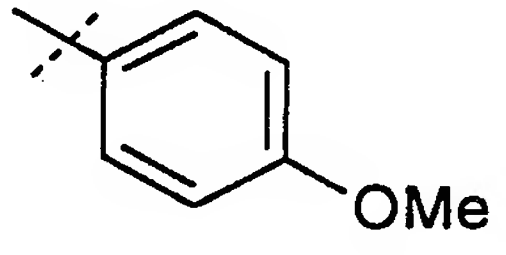
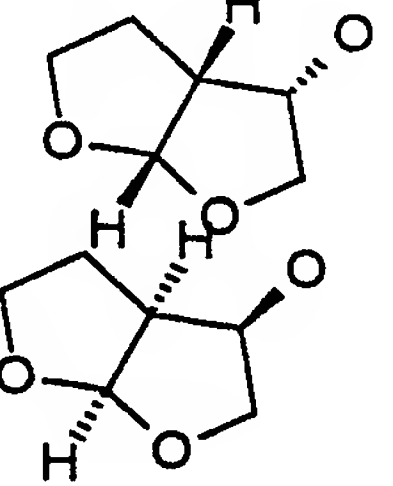
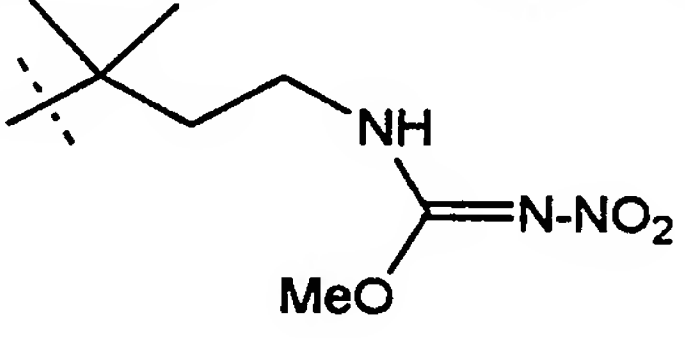
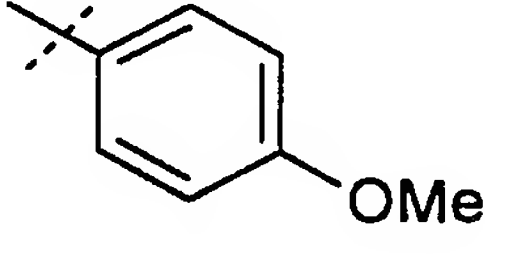
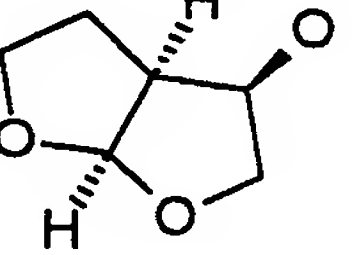
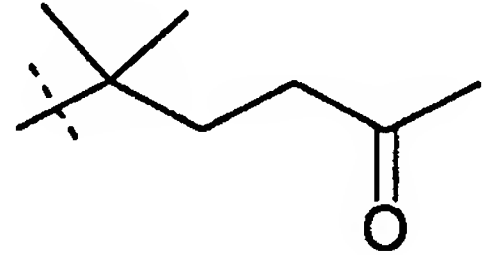
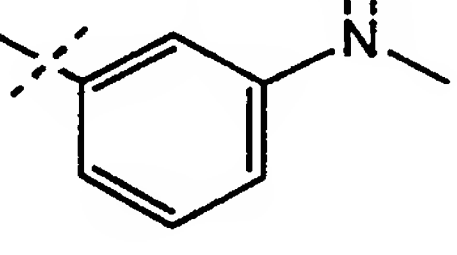
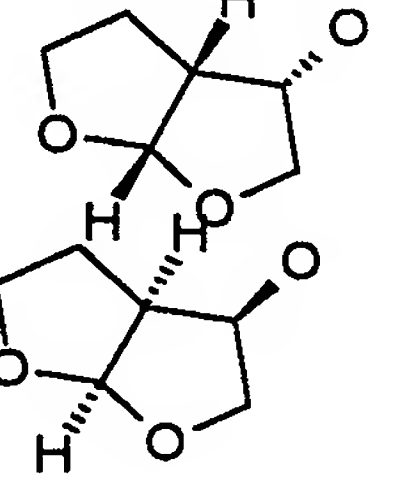
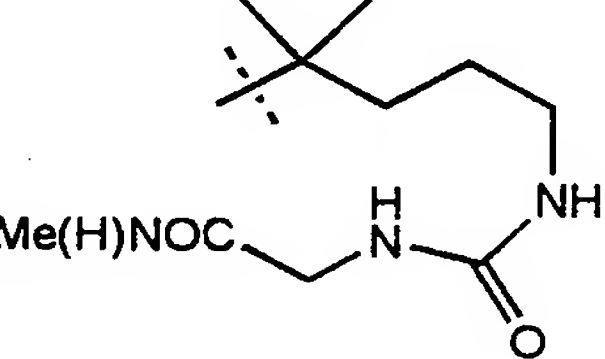
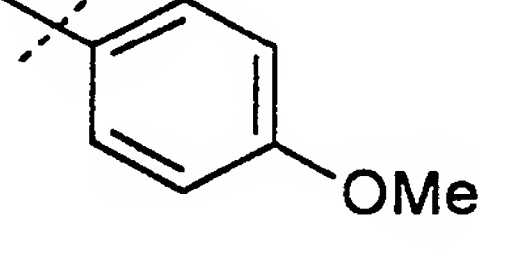
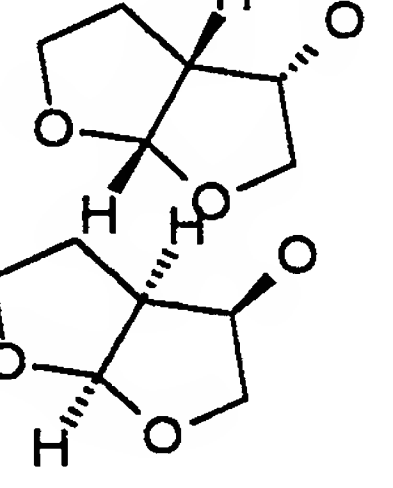
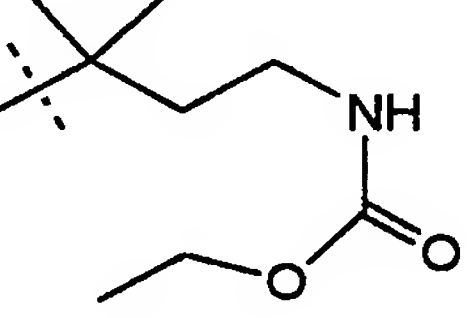
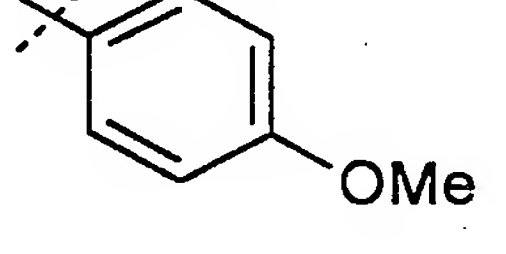
Compound	R'	R''	E
29			
30			
31			
32			
33			
35			
36			
37			

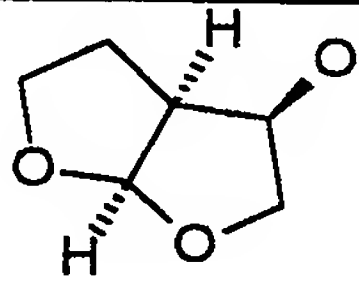
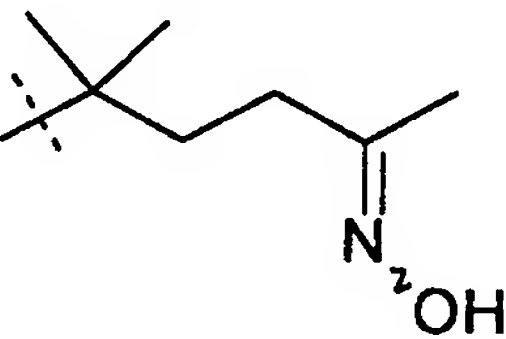
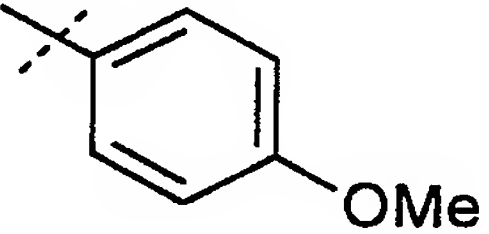
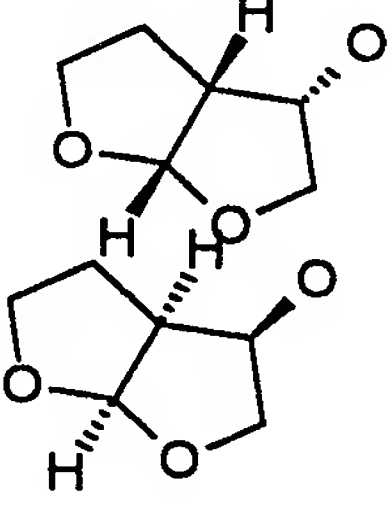
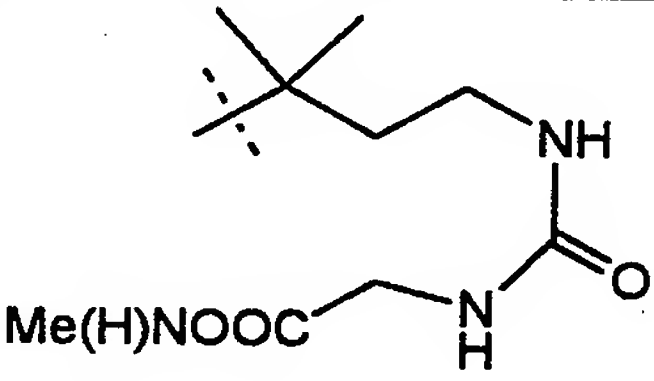
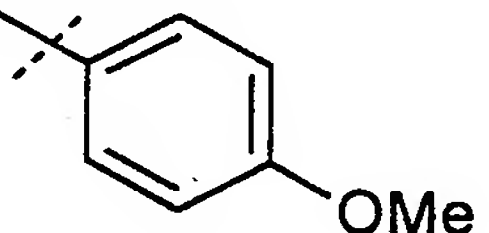
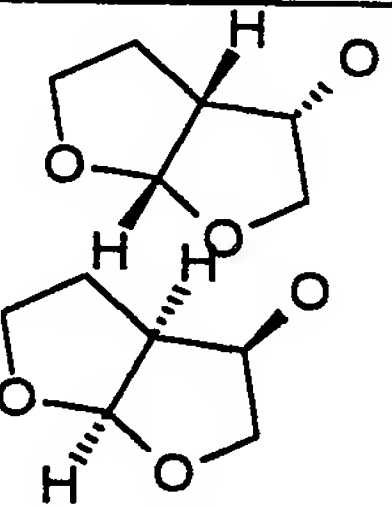
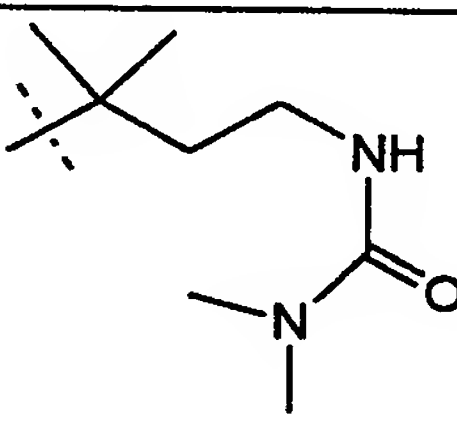
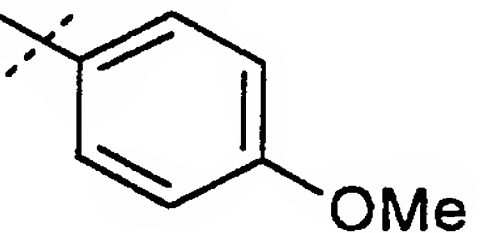
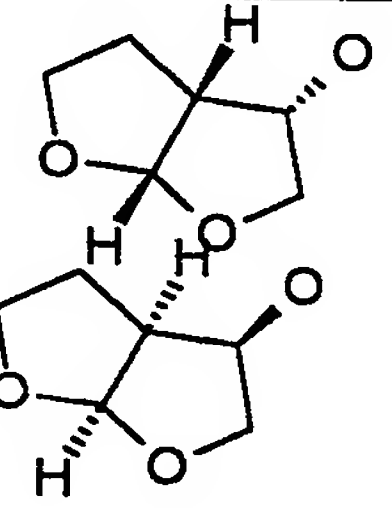
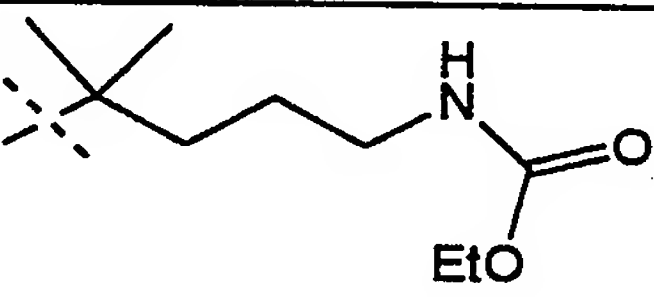
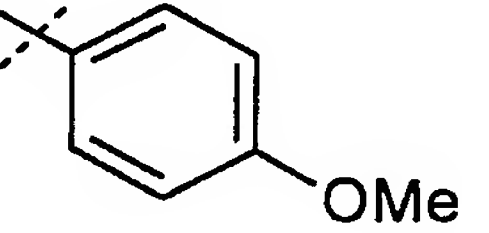
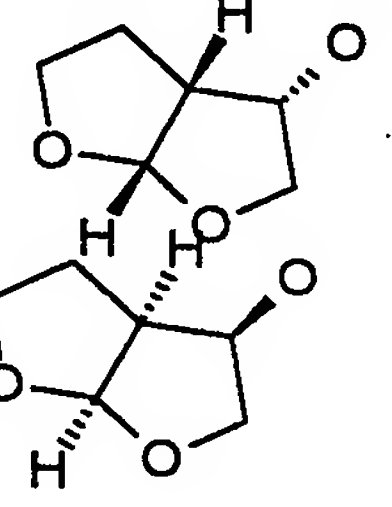
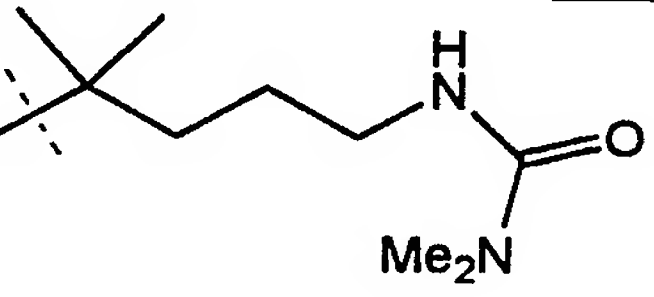
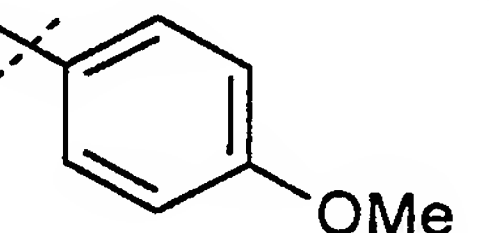
Compound	R'	R''	E
38			
39			
40			
41			
42			
43			
44			
124			

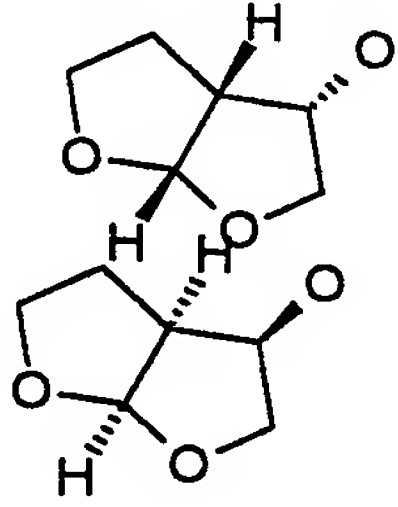
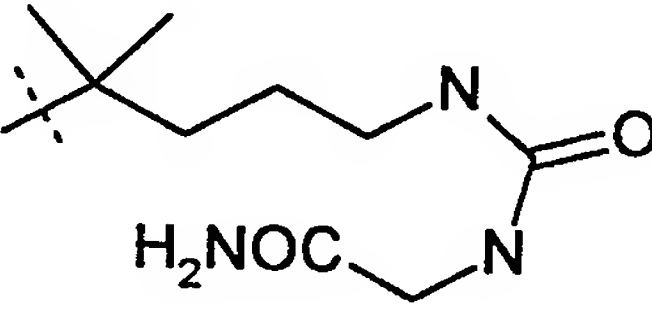
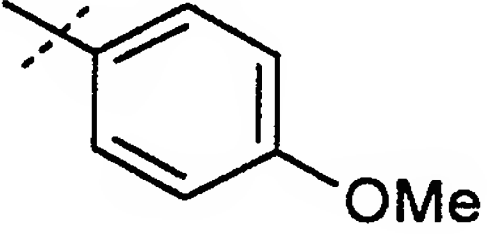
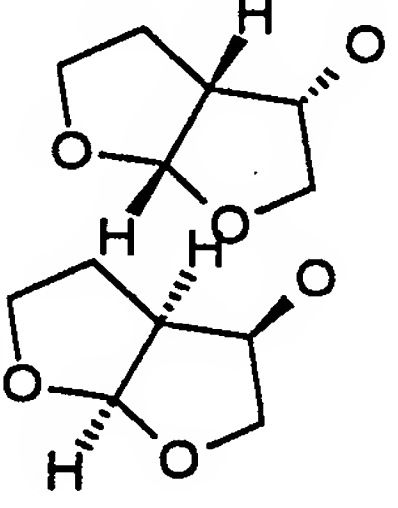
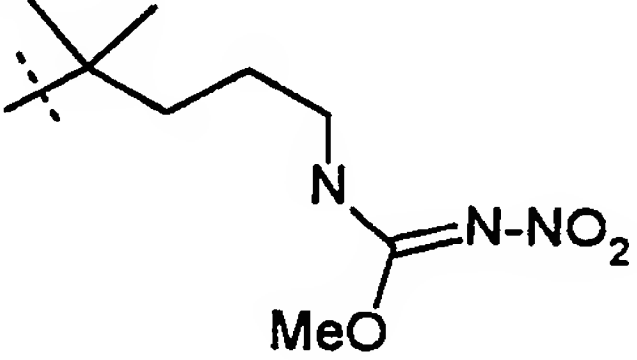
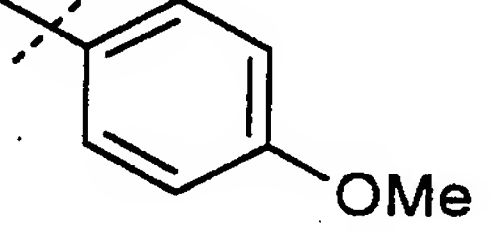
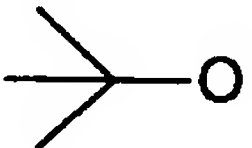
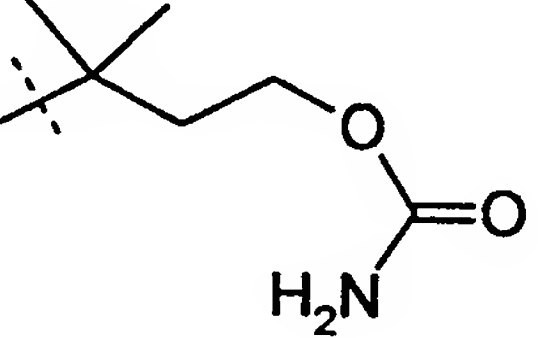
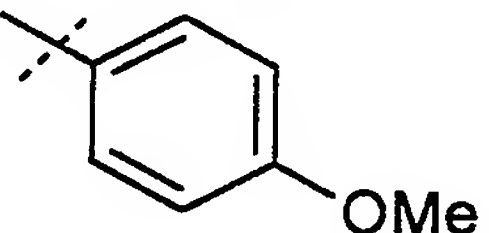
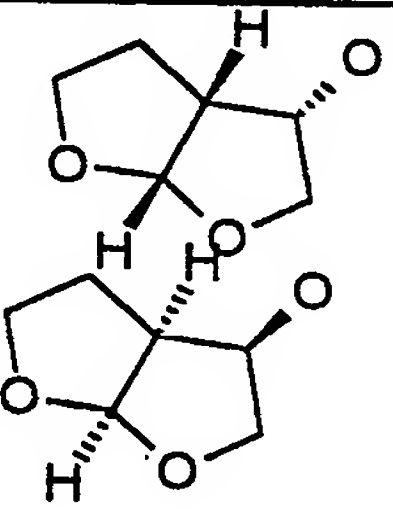
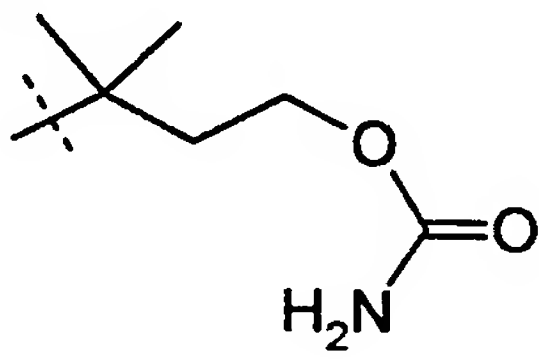
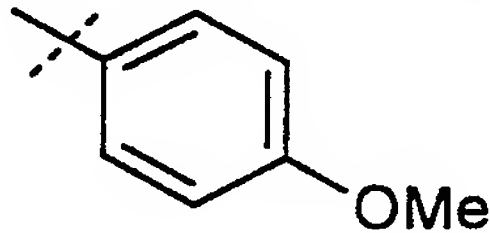
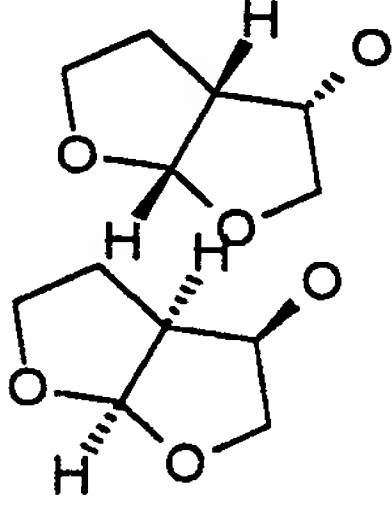
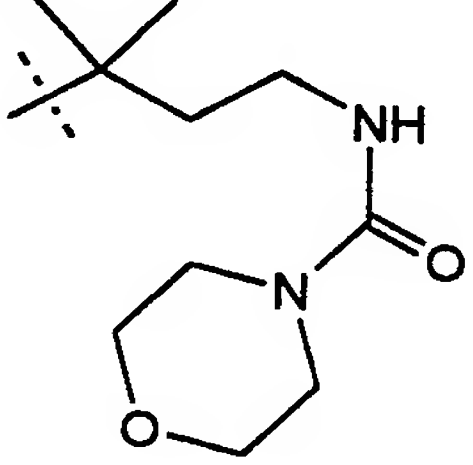
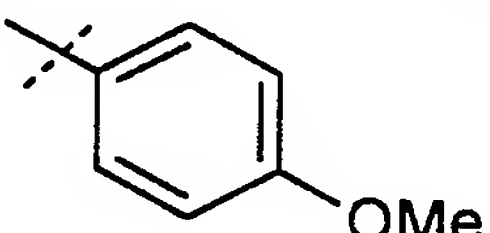
Compound	R'	R''	E
125			
127			
203			
205			
206			
207			
208			
209			
210			

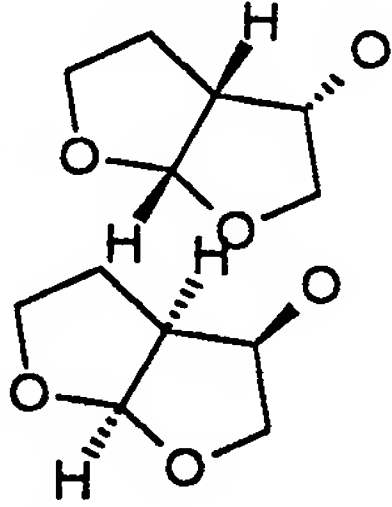
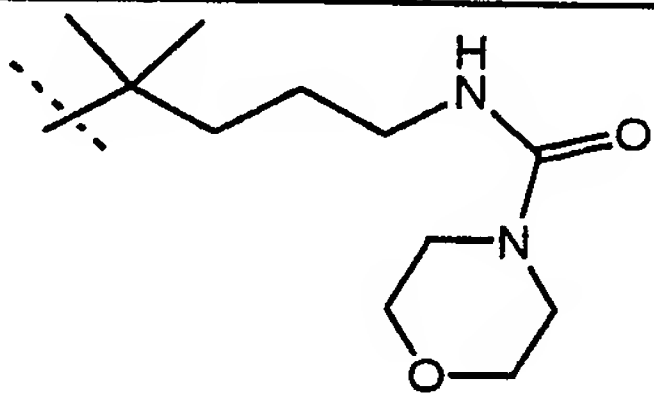
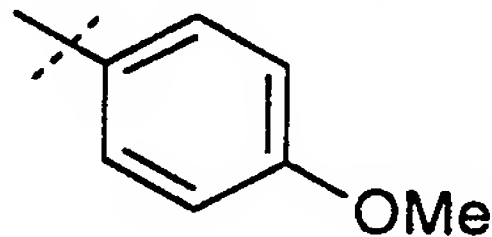
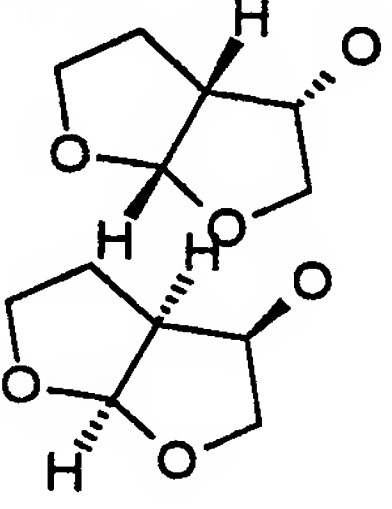
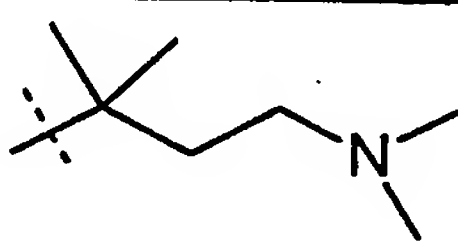
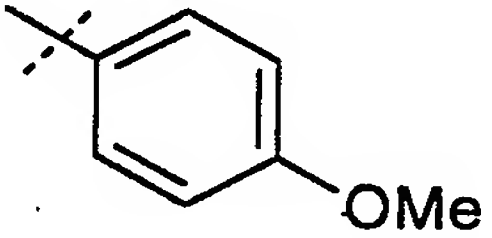
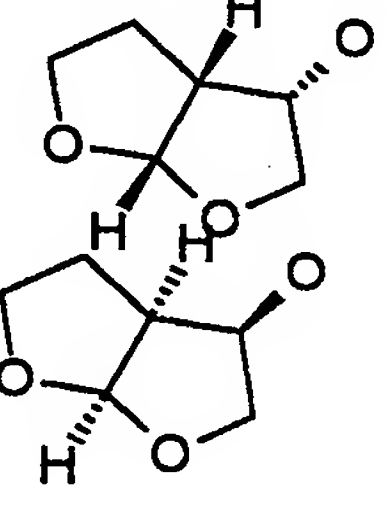
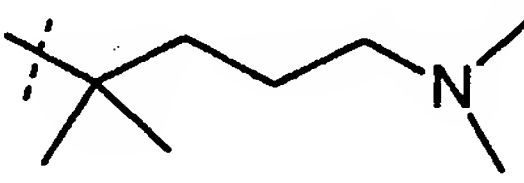
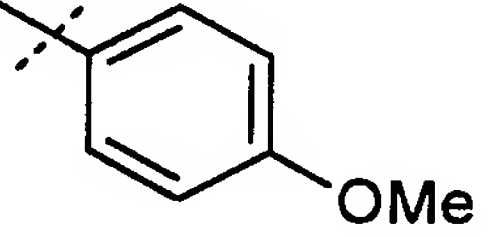
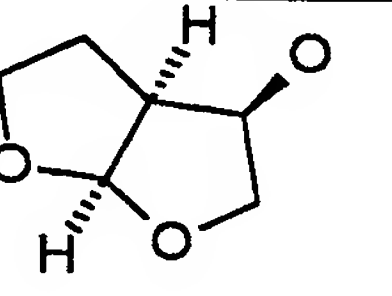
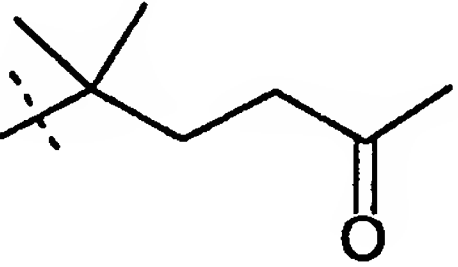
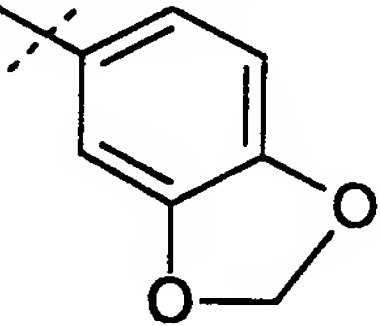
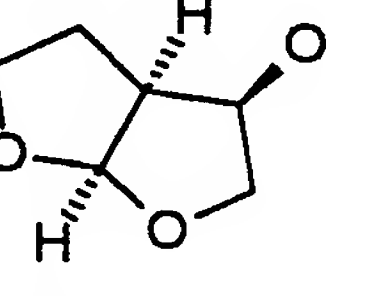
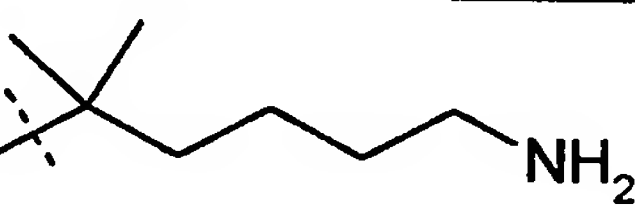
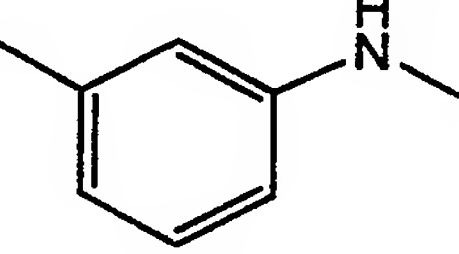
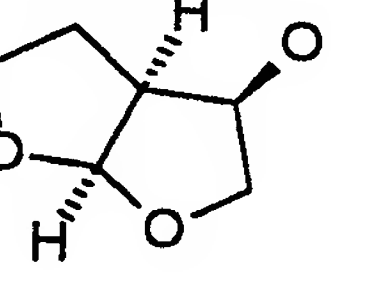
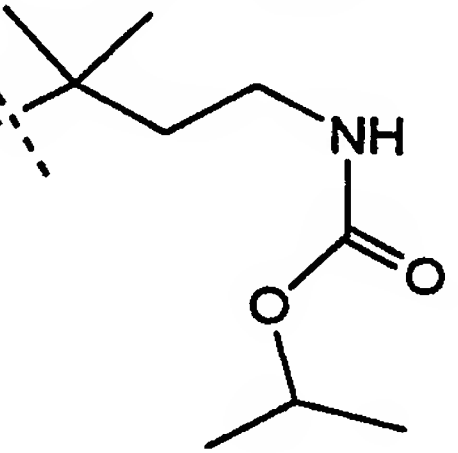
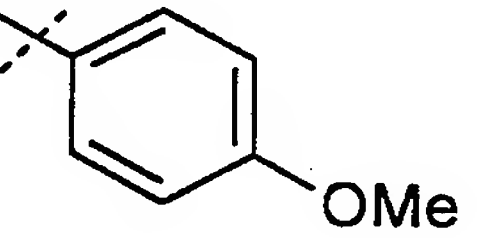
Compound	R'	R''	E
211			
212			
213			
214			
215			

Compound	R'	R''	E
216			
217			
218			
219			
220			

Compound	R'	R''	E
221			
222			
223			
224			
225			
226			

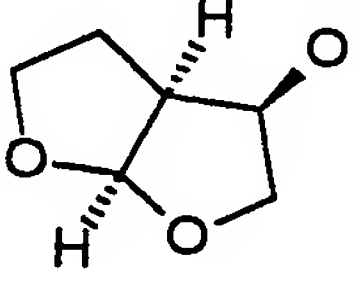
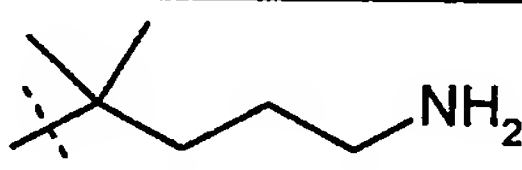
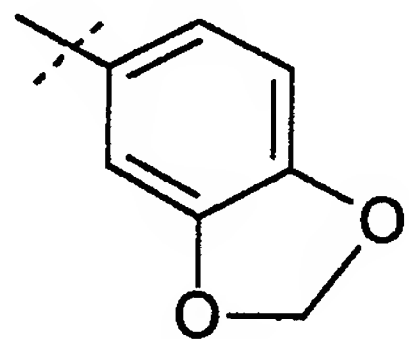
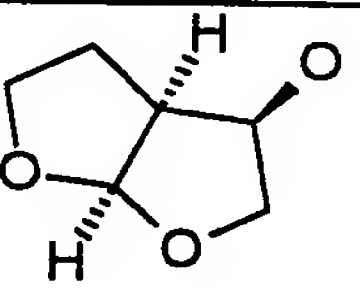

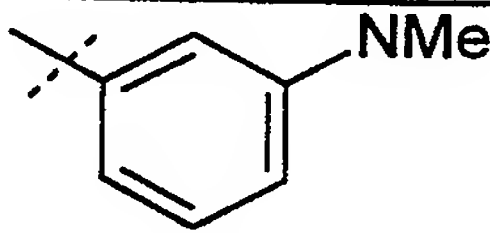
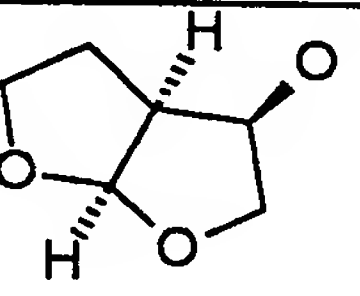
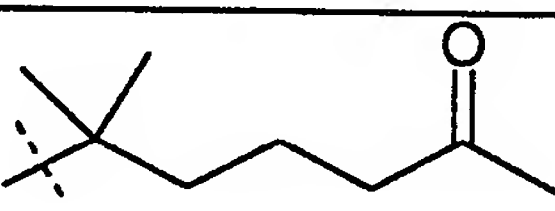
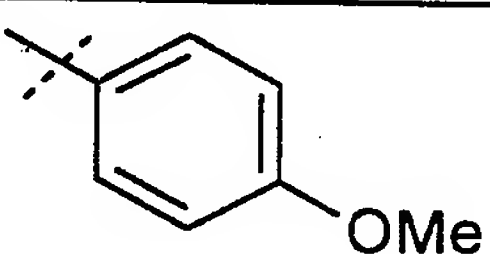
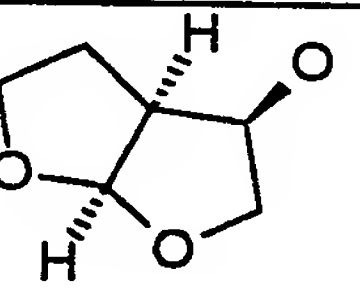
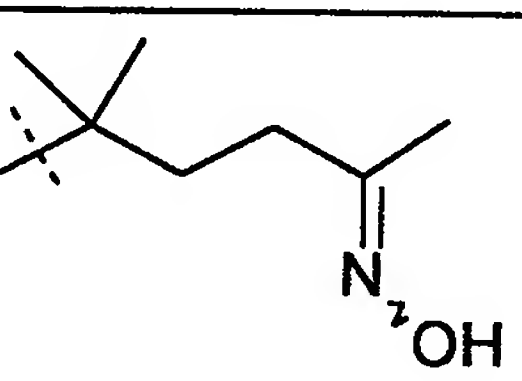
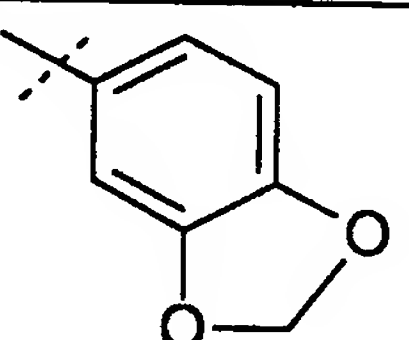
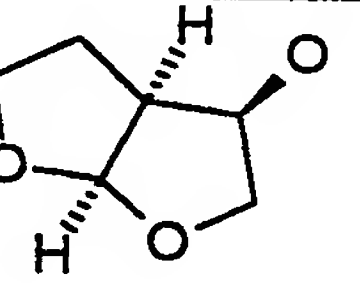
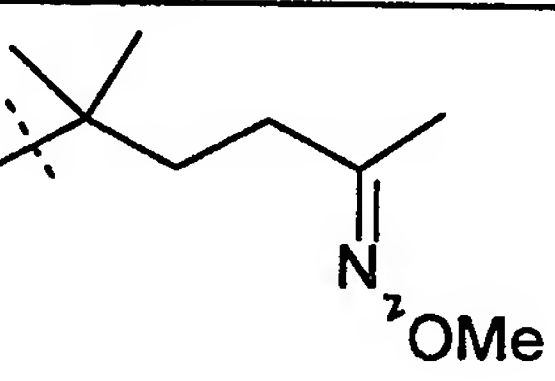
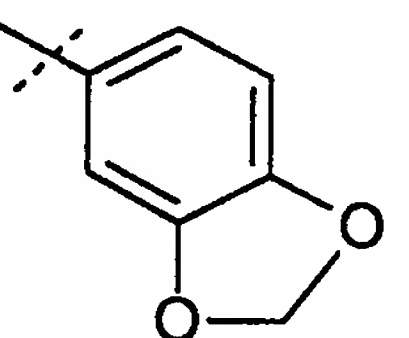
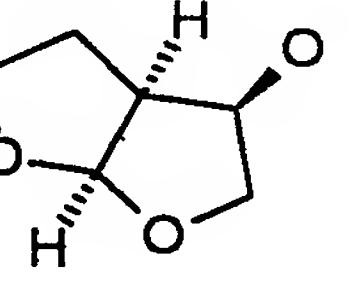
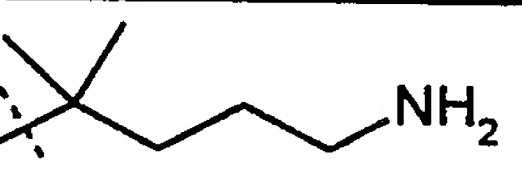
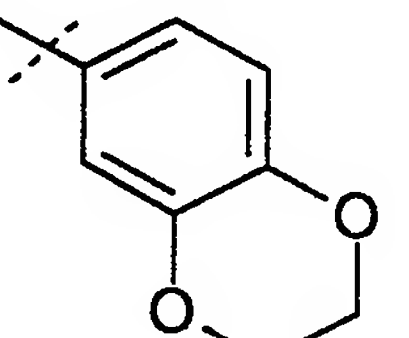
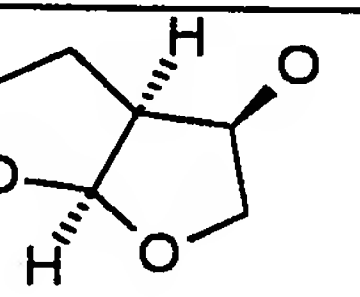
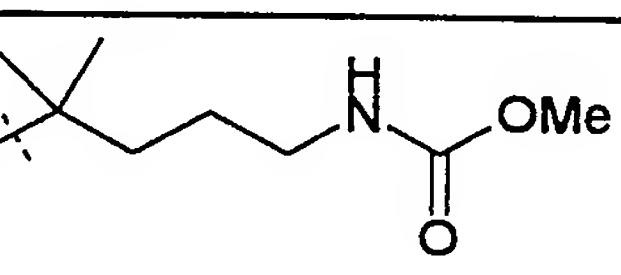
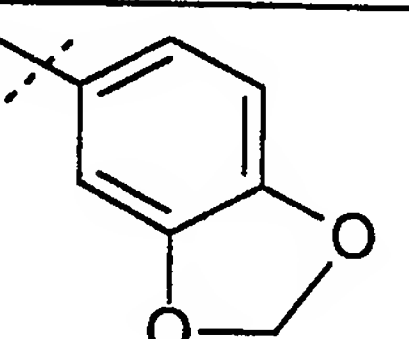
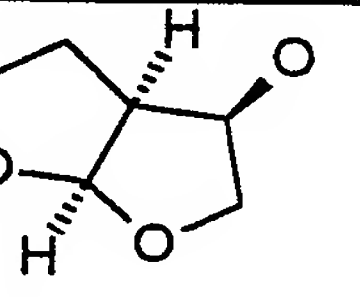
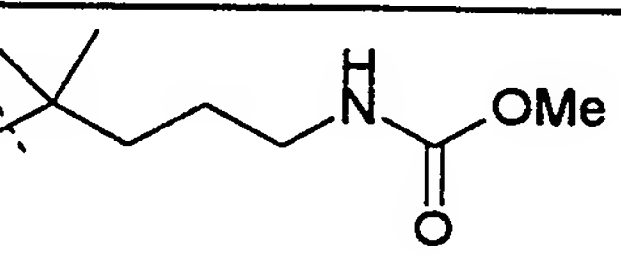
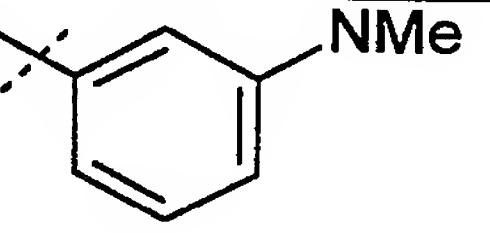
Compound	R'	R''	E
227			
228			
229			
230			
231			

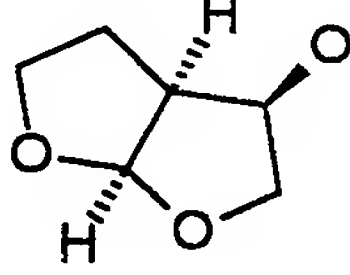
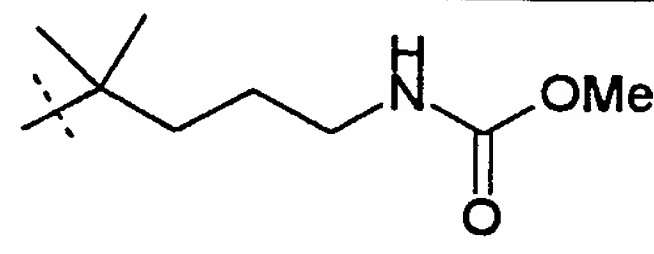
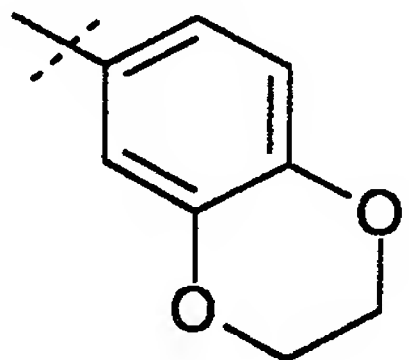
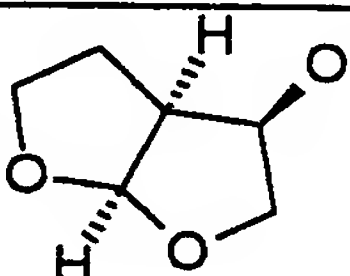
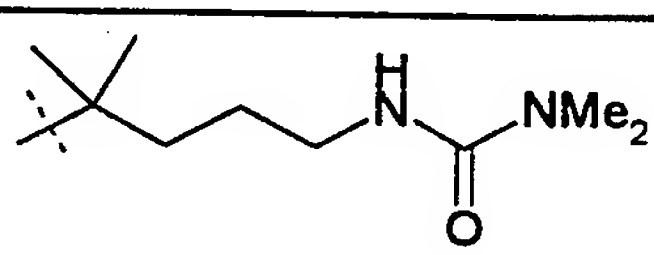
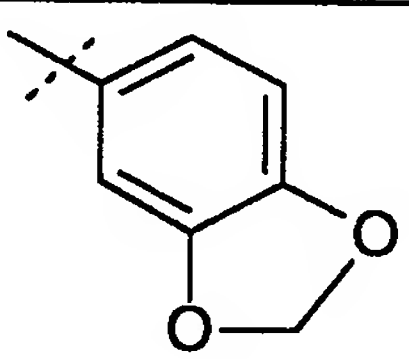
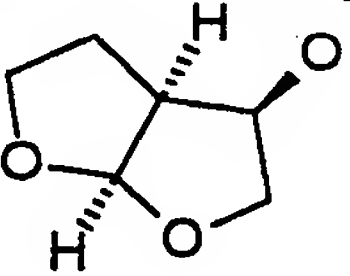
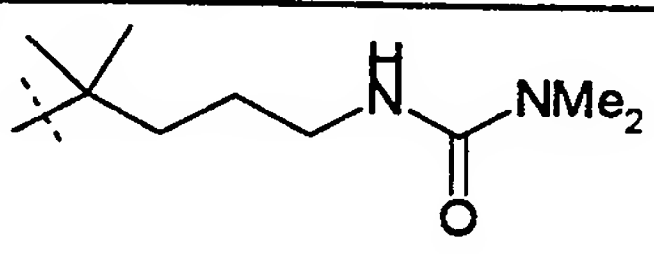
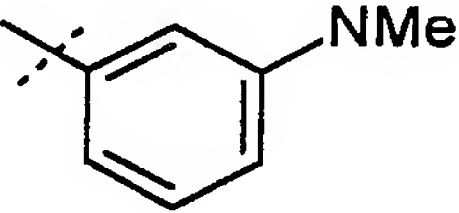
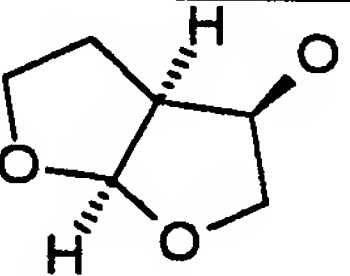
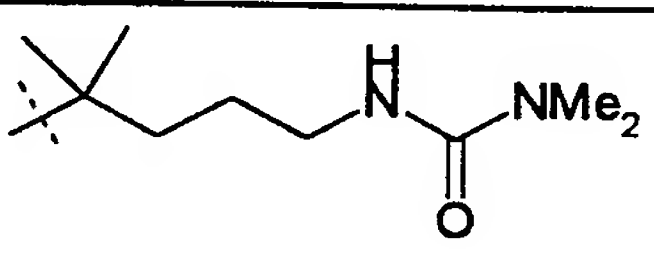
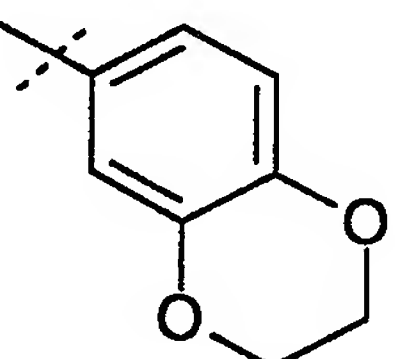
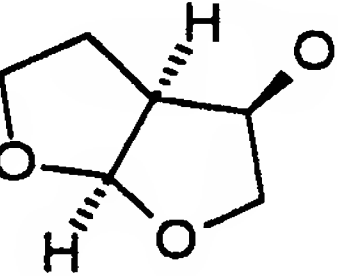
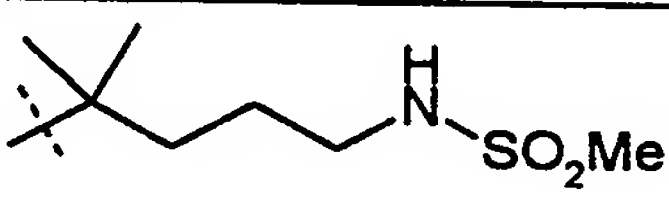
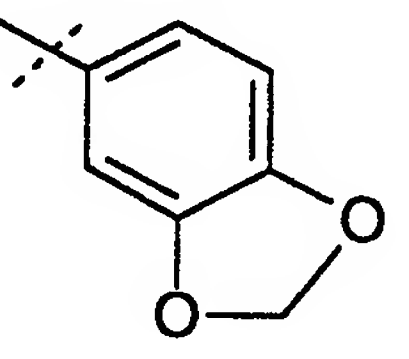
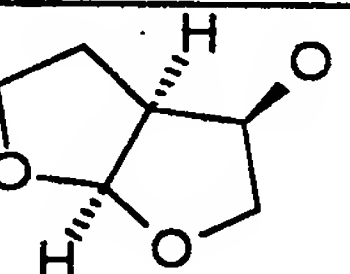
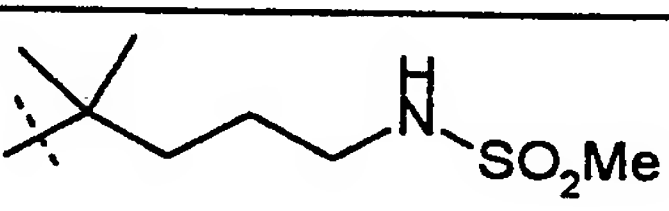
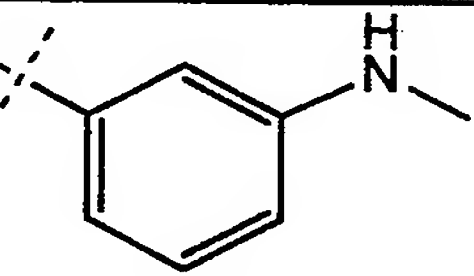
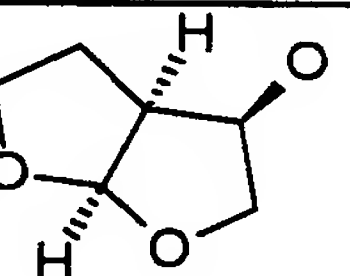
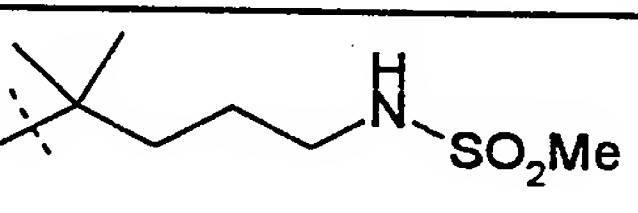
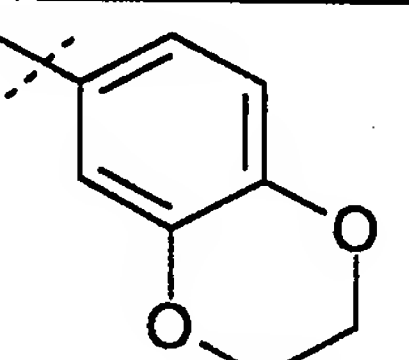
Compound	R'	R''	E
232		 <chem>CC(C)(C)CCN(C(=O)OCCN(C(=O)OCC(C)(C)C)C(C)(C)C)C(C)(C)C</chem>	 <chem>COc1ccc(cc1)C(C)(C)C</chem>
233		 <chem>CC(C)(C)CCN(C(=O)OCCN(C(=O)OCC(C)(C)C)C(C)(C)C)C(C)(C)C</chem>	 <chem>COc1ccc(cc1)C(C)(C)C</chem>
234		 <chem>CC(C)(C)CCN(C(=O)OCCN(C(=O)OCC(C)(C)C)C(C)(C)C)C(C)(C)C</chem>	 <chem>COc1ccc(cc1)C(C)(C)C</chem>
235		 <chem>CC(C)(C)CCN(C(=O)OCCN(C(=O)OCC(C)(C)C)C(C)(C)C)C(C)(C)C</chem>	 <chem>COc1ccc(cc1)C(C)(C)C</chem>
236		 <chem>CC(C)(C)CCN(C(=O)OCCN(C(=O)OCC(C)(C)C)C(C)(C)C)C(C)(C)C</chem>	 <chem>COc1ccc(cc1)C(C)(C)C</chem>

Compound	R'	R''	E
237			
238			
239			
240			
241			
242			

Compound	R'	R''	E
243			
244			
245			
246			
247			
248			
249			

Compound	R'	R''	E
250			
251			
252			
253			
254			
255			
256			

Compound	R'	R''	E
257			
258			
259			
260			
261			
262			
263			
264			

Compound	R'	R''	E
265			
266			
267			
268			
269			
270			
271			

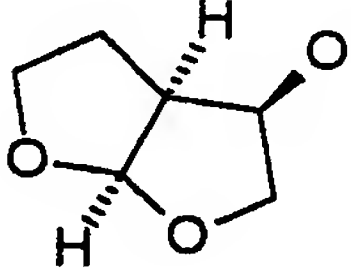
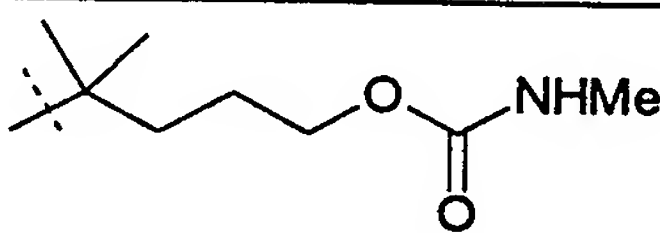
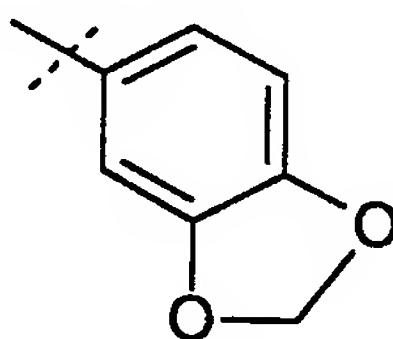
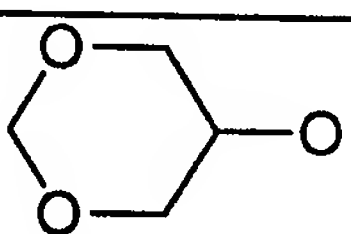
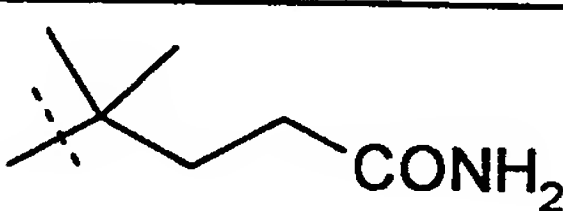
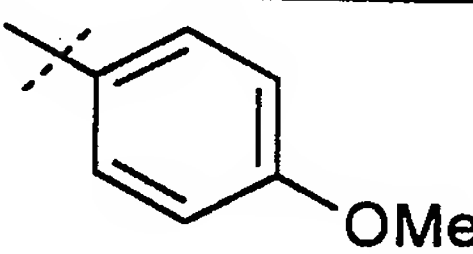
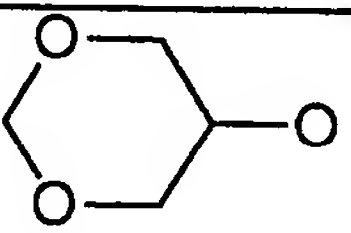
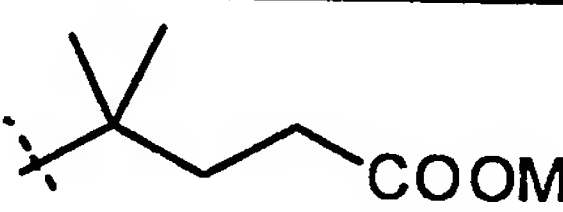
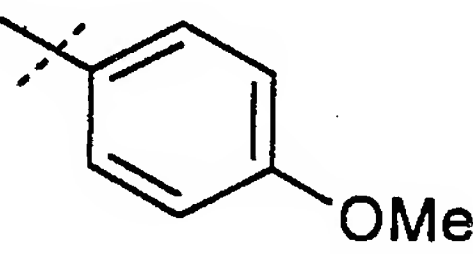

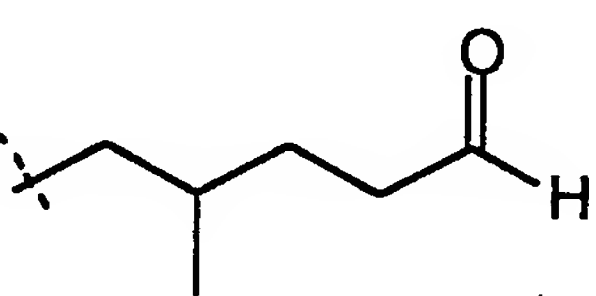
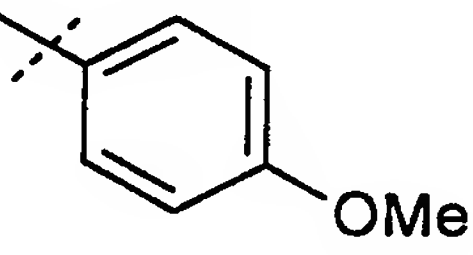
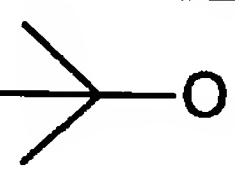
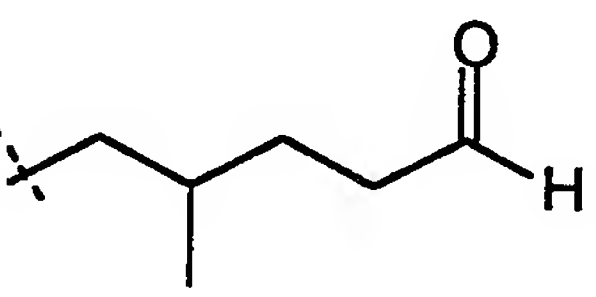
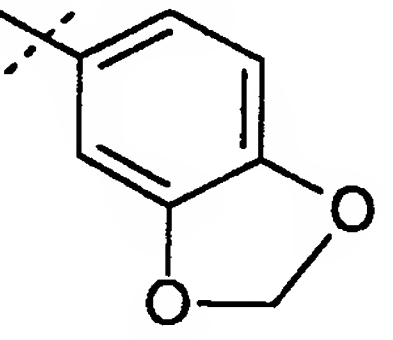
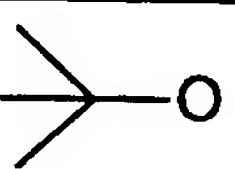
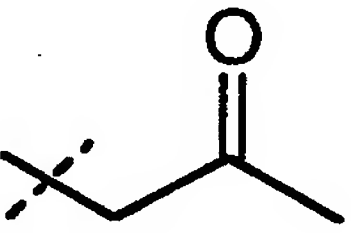
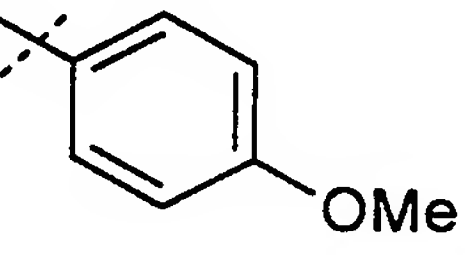
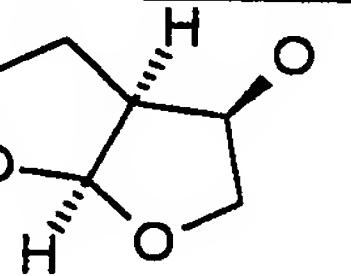
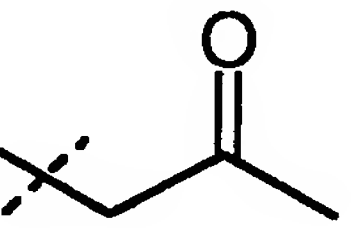
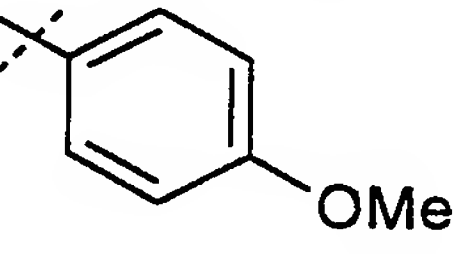
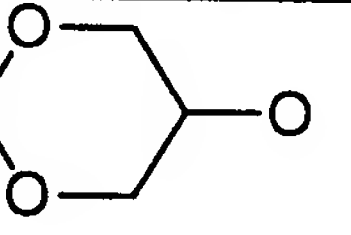
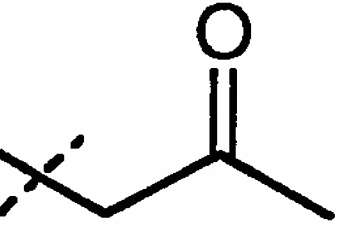
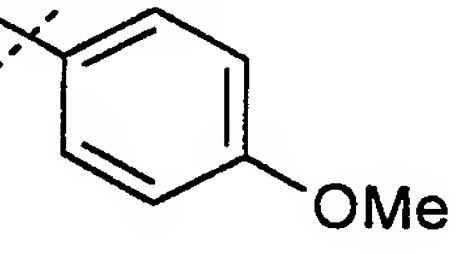
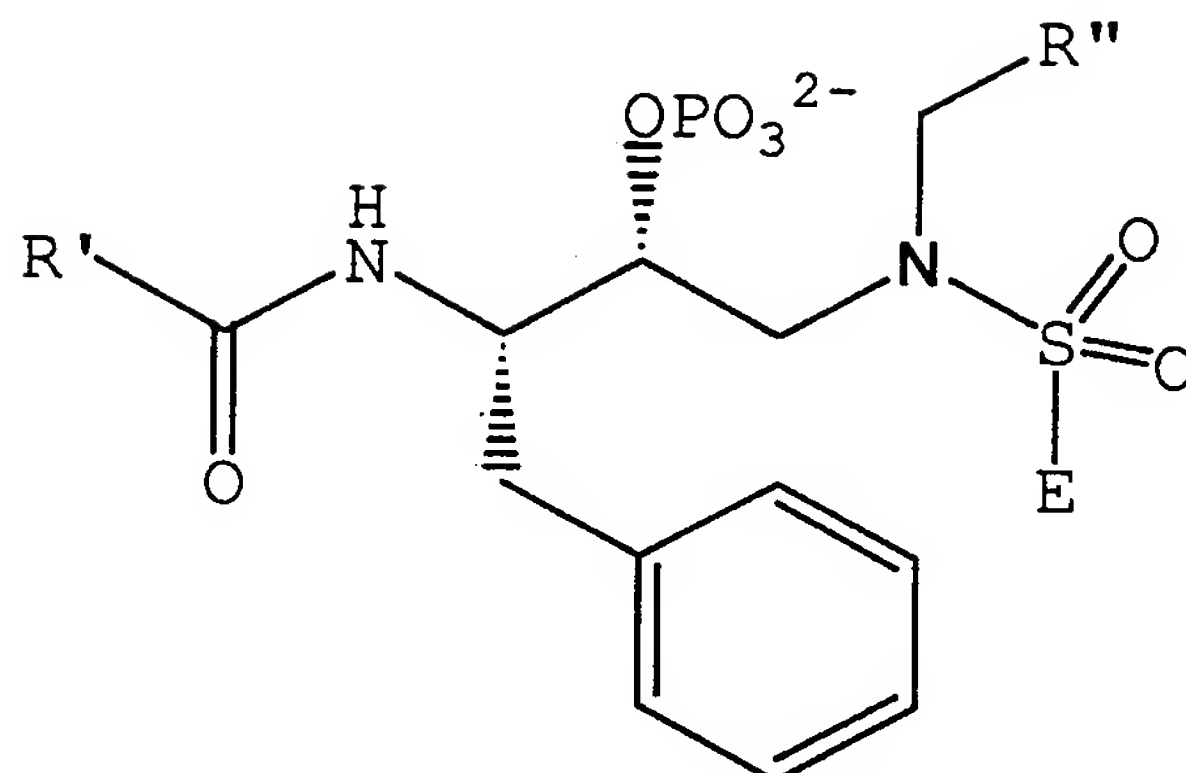
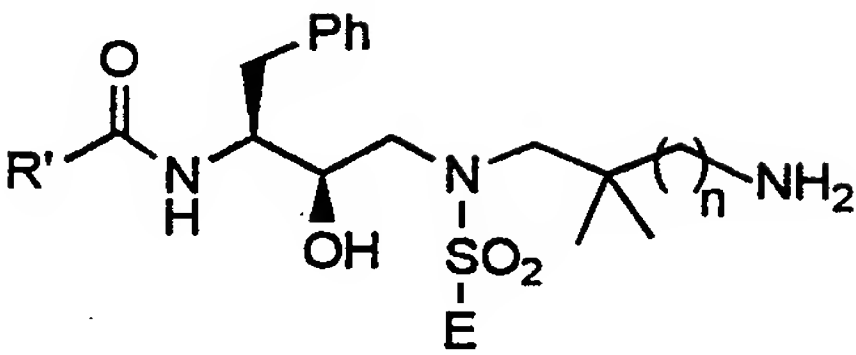
Compound	R'	R''	E
272			
273			
274			
277			
278			
279			
280			
281			

Table 2



Compound	R'	R''	E
100			
101			
47			

Table 3



Compound	R'	n	E
282		2	
283		5	
284		2	
285		2	
286		2	
287		3	
288		3	
289		4	

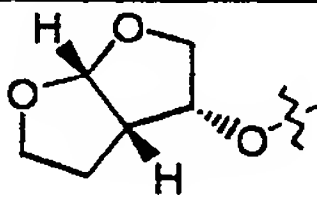

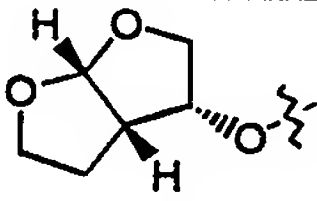
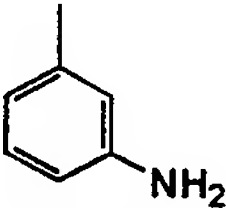
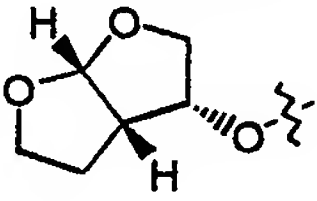
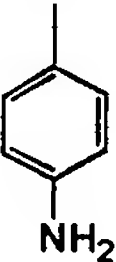
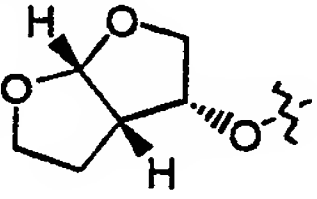
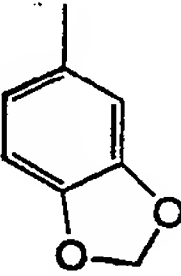
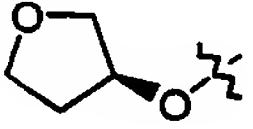
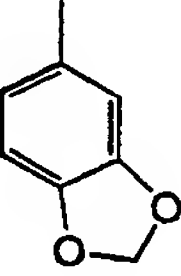
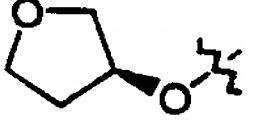
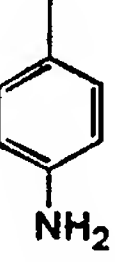
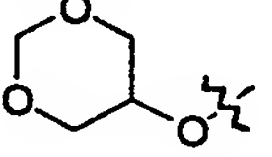
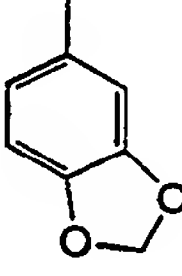
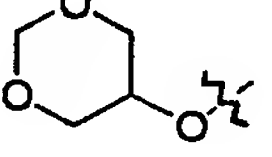
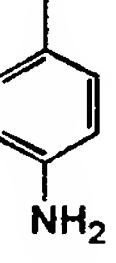
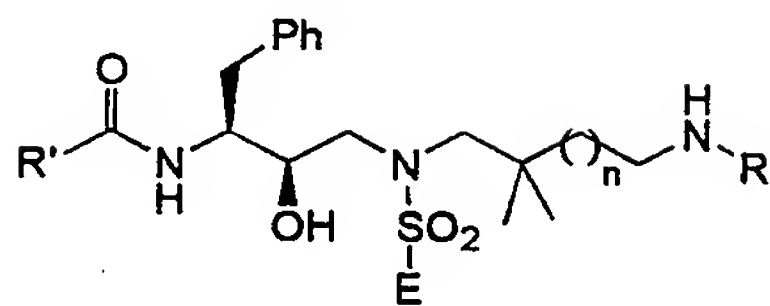
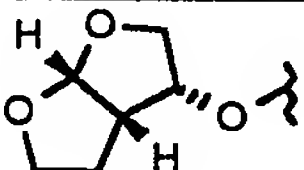
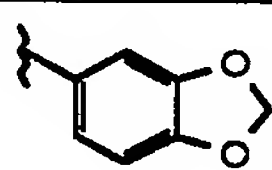
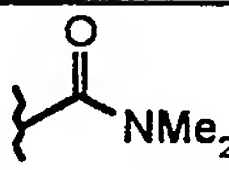
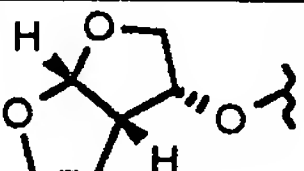
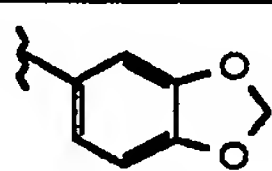
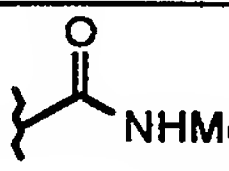
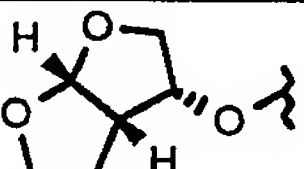
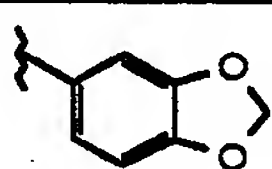
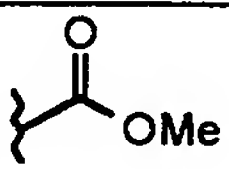
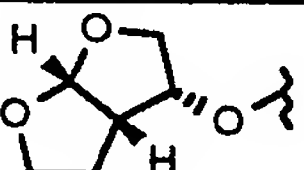
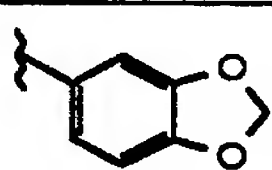
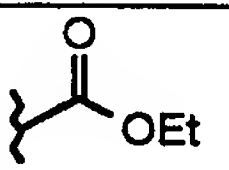
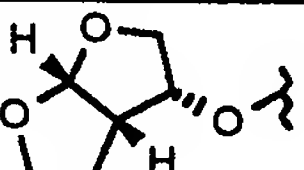
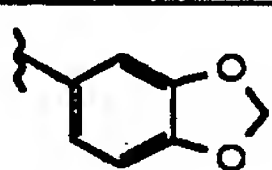
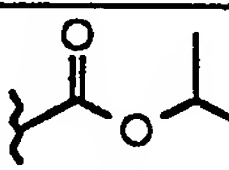
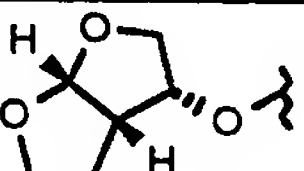
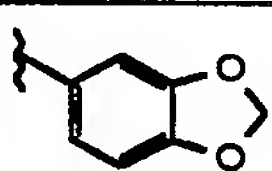
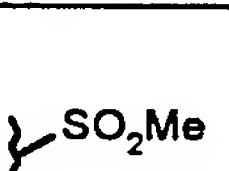
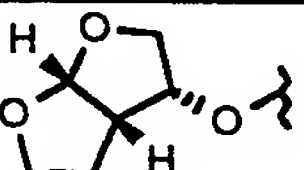
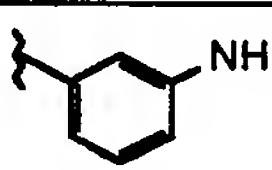
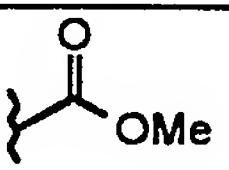
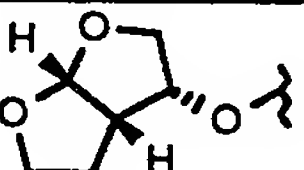
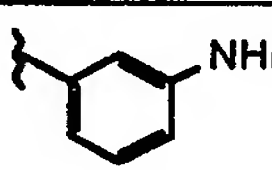

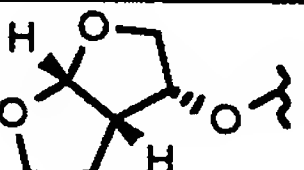
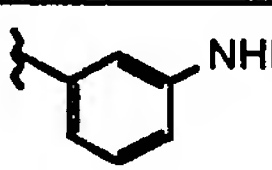
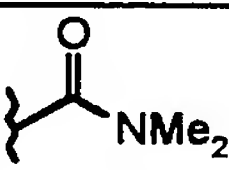
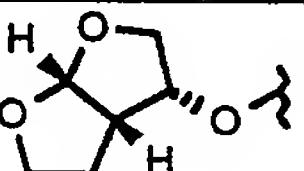
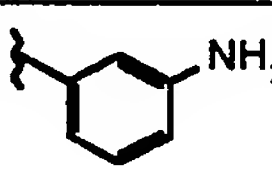
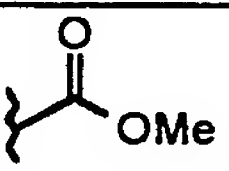
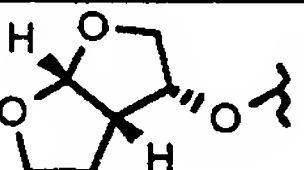
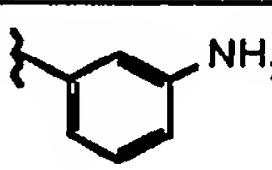
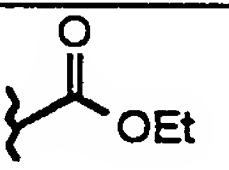
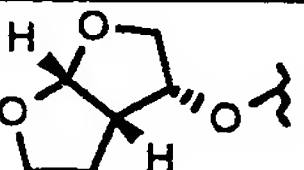
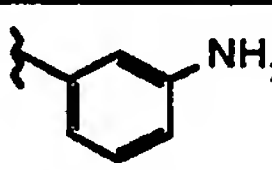
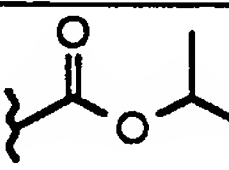
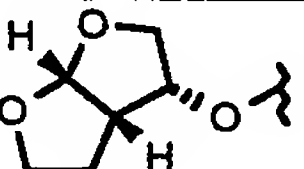
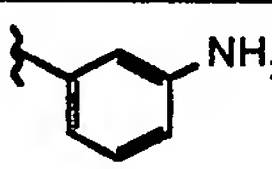
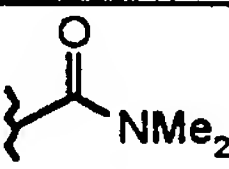
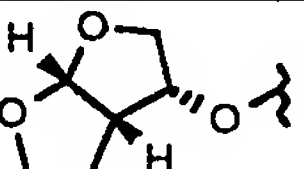
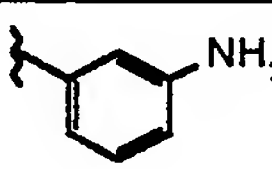
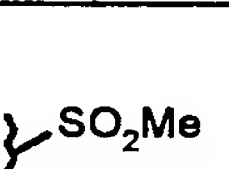
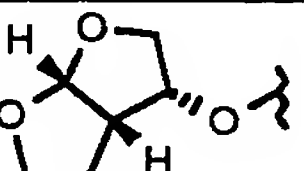
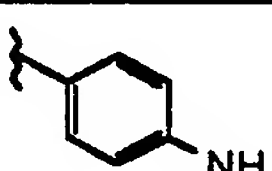
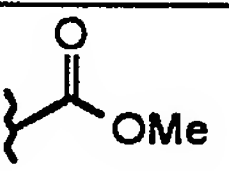
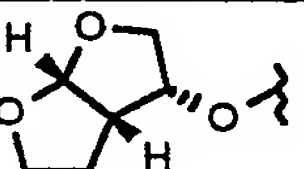
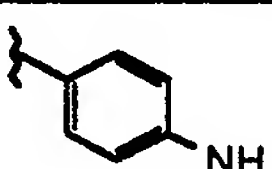
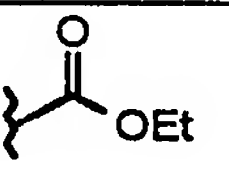
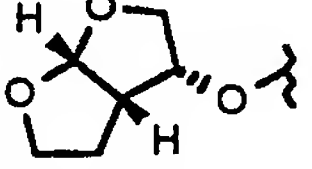
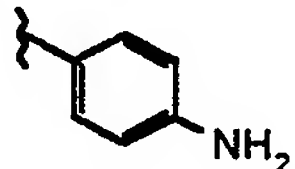
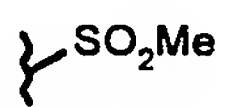
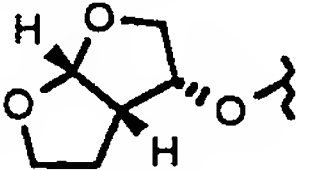
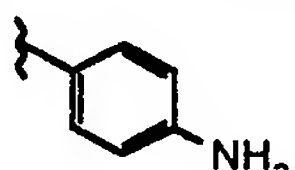
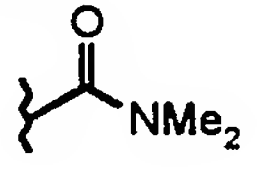
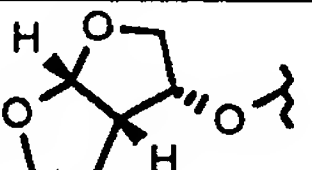
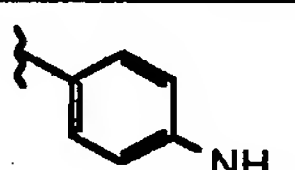
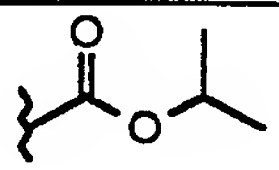
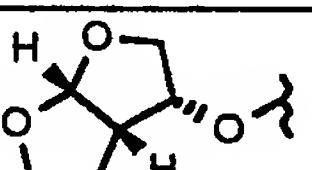
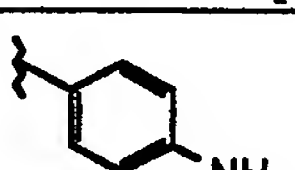
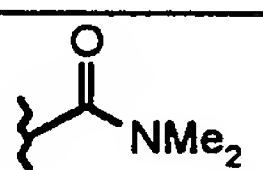
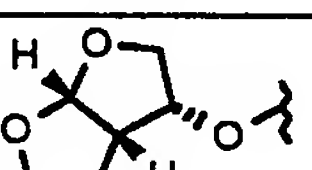
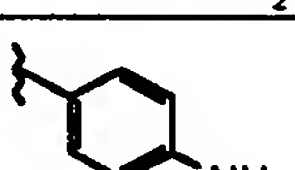
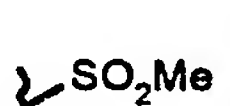
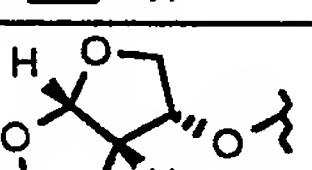
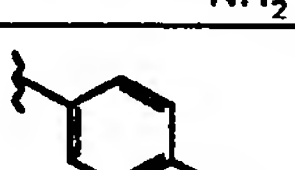
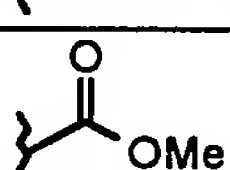
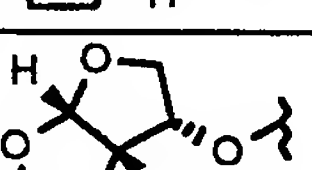

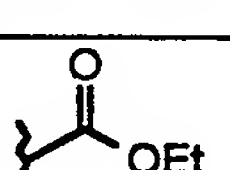
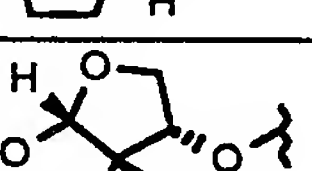
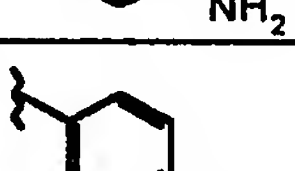
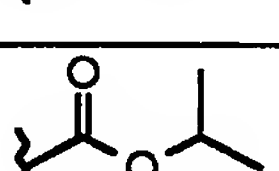
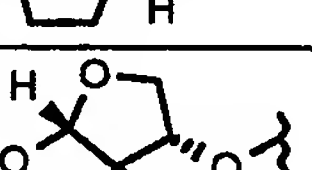
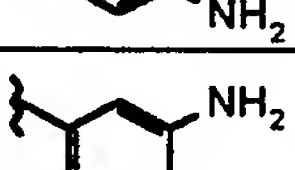
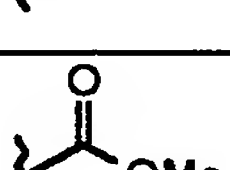
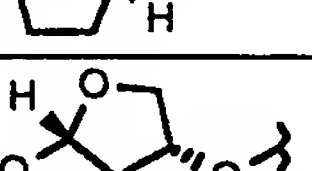
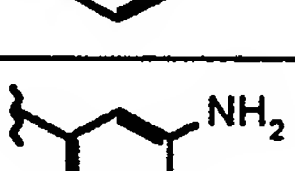
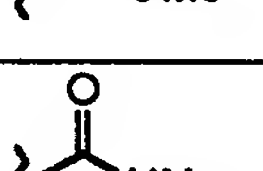
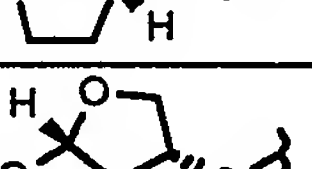
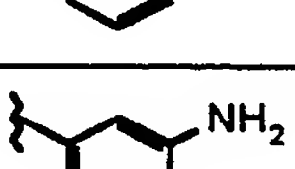

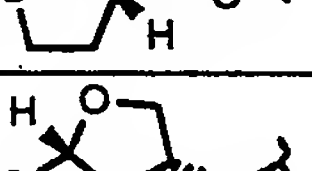
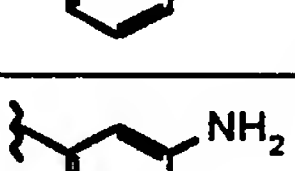
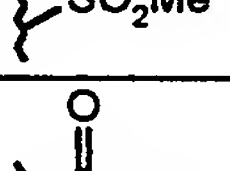
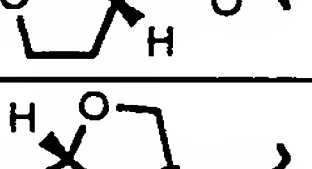
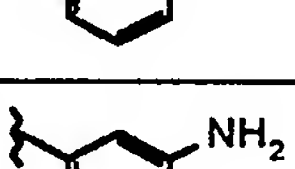
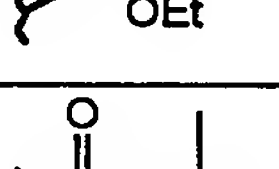
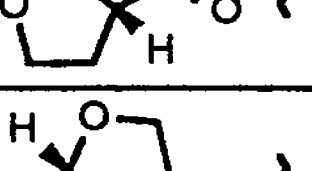
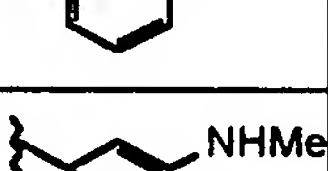
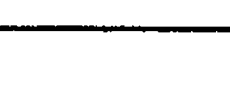
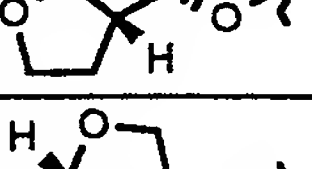
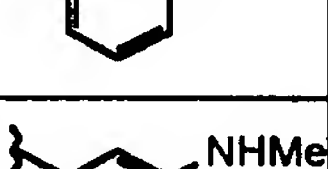
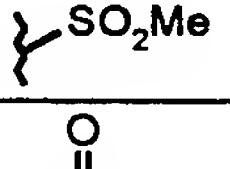
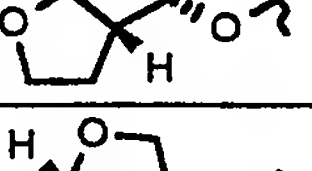
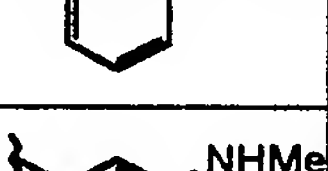
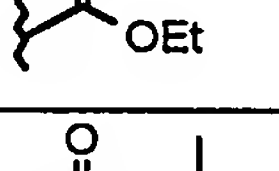
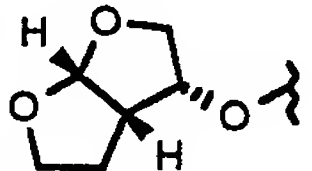
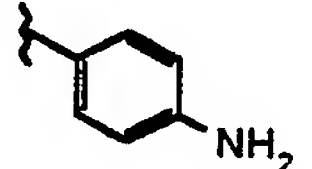
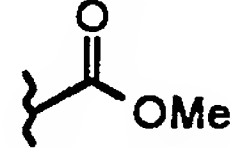
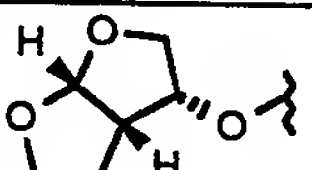
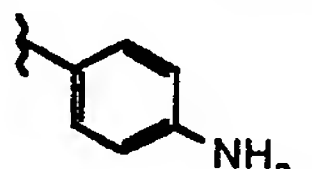
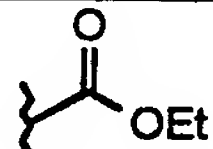
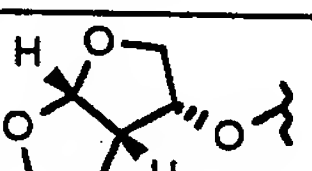
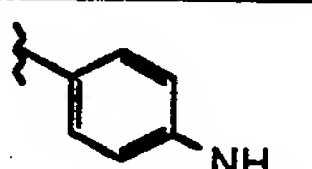
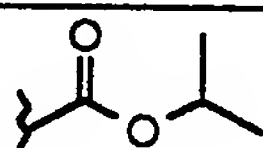
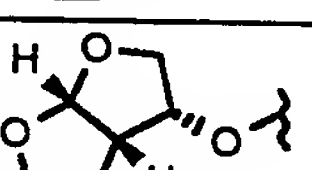
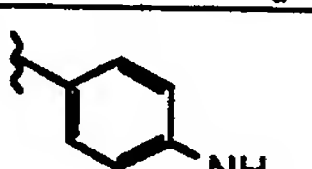
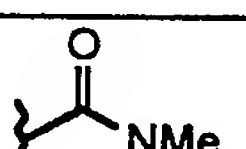
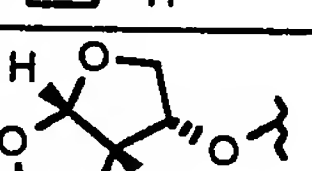
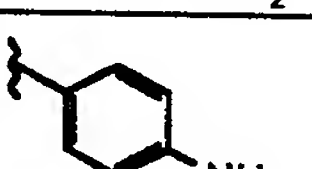
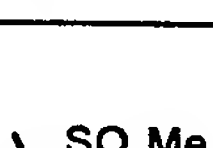
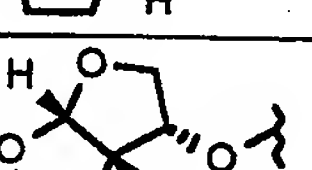
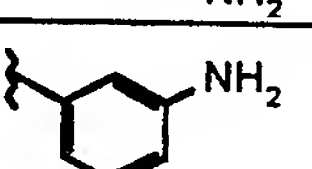
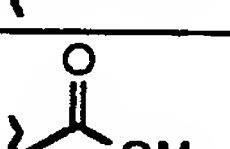
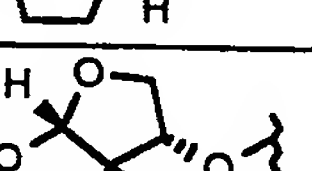

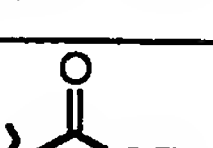
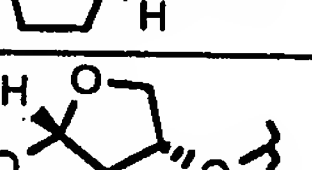
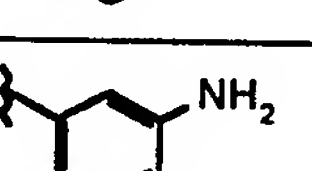
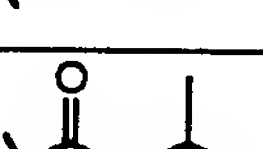
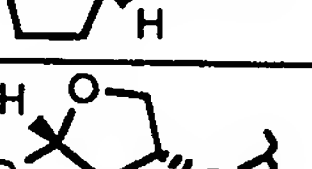
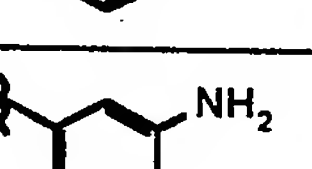

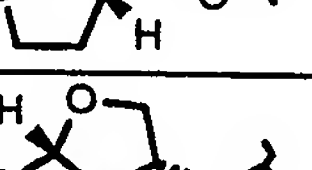
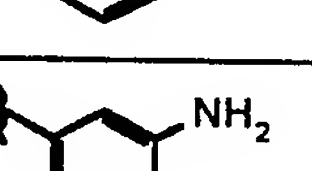
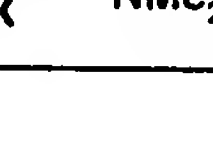
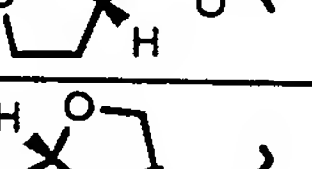
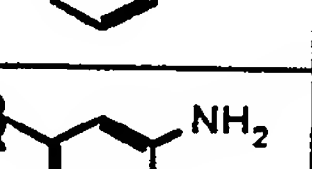
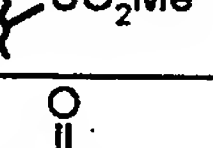
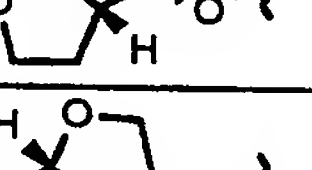
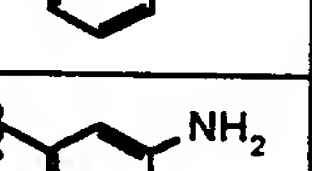
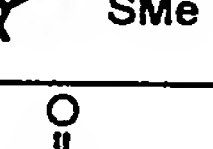
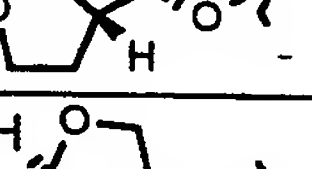

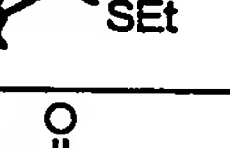
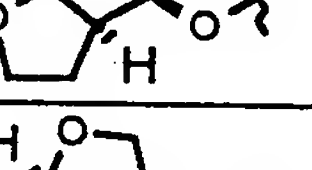
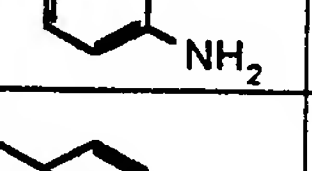
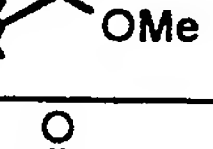
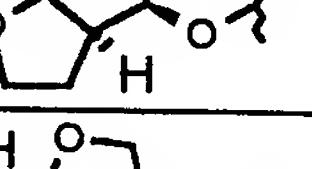
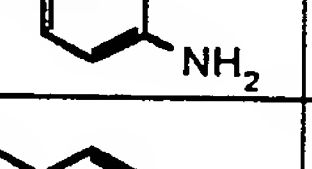
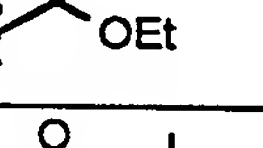
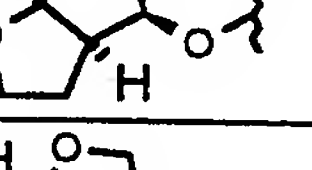
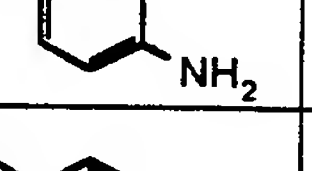
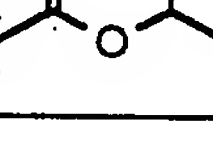
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298		4	

Table 4

Compound	R'	E	n	R
299			3	
300			3	
301			3	
302			2	
303			2	
304			2	
305			2	
306			2	
307			1	
308			1	
309			1	
310			1	

311			1	
312			1	
313			4	
314			4	
315			4	
316			4	
317			1	
318			1	
319			1	
320			1	
321			1	
322			1	
323			1	
324			1	
325			1	
326			1	

327			1	
328			1	
329			1	
330			2	
331			2	
332			2	
333			2	
334			2	
335			2	
336			2	
337			2	
338			2	
339			2	
340			3	
341			3	
342			3	

343			3	
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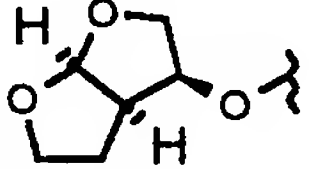
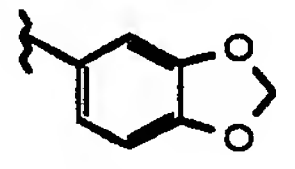
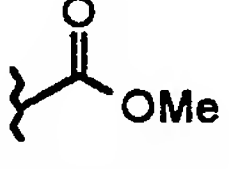
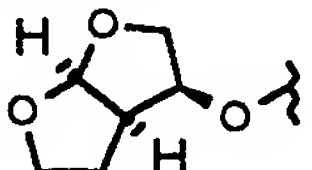
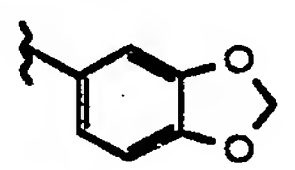
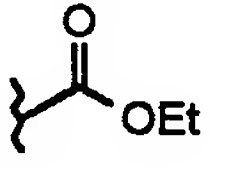
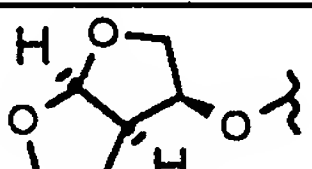
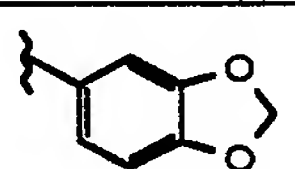
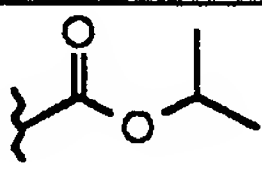
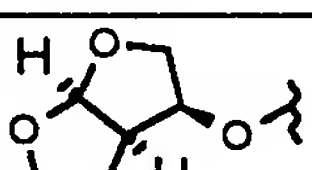
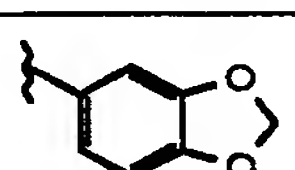
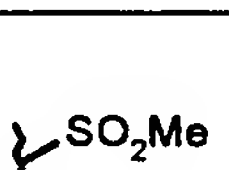
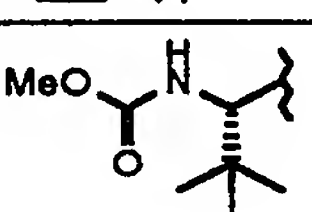
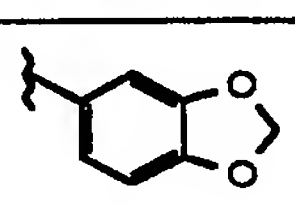
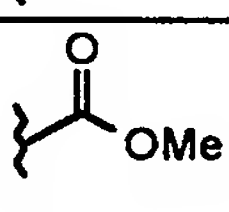
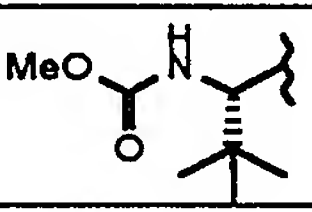
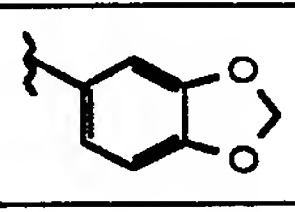
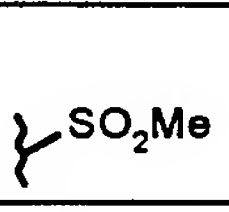
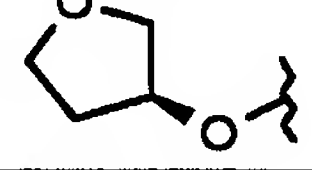
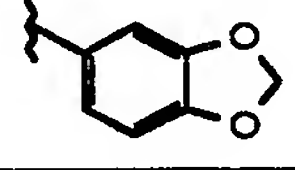
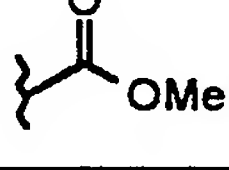
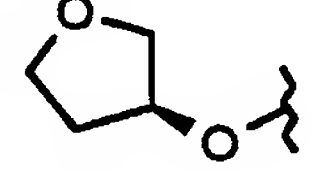
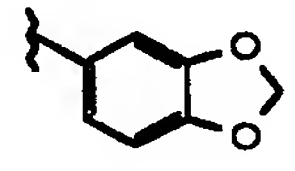
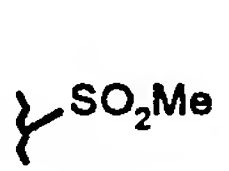
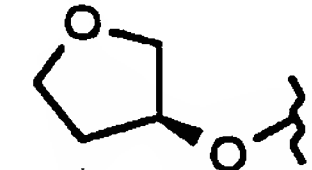
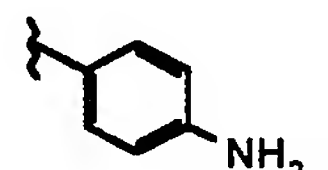
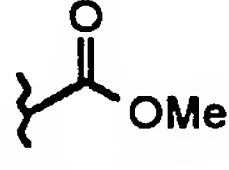
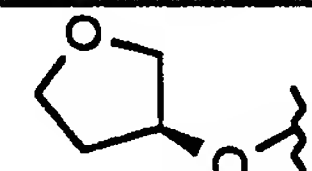
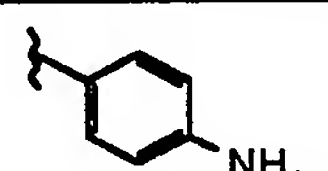
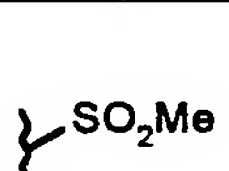
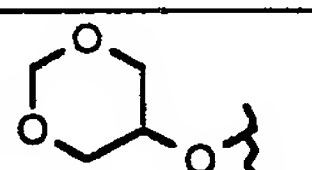
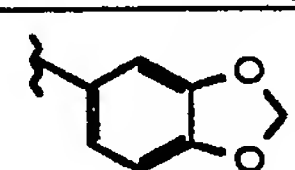
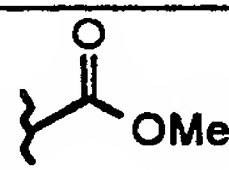
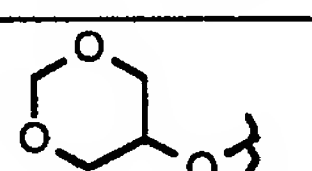
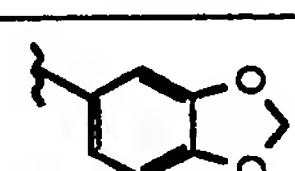
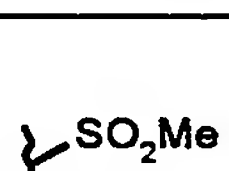
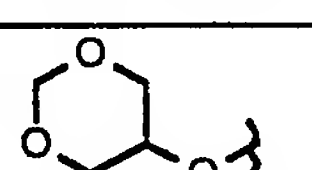
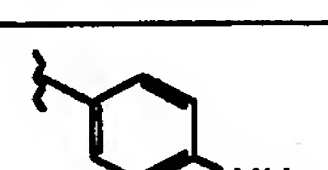
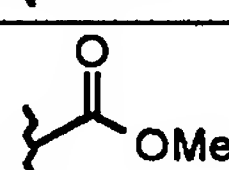
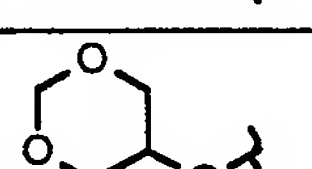

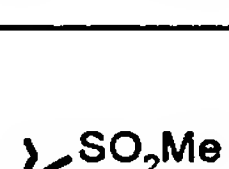
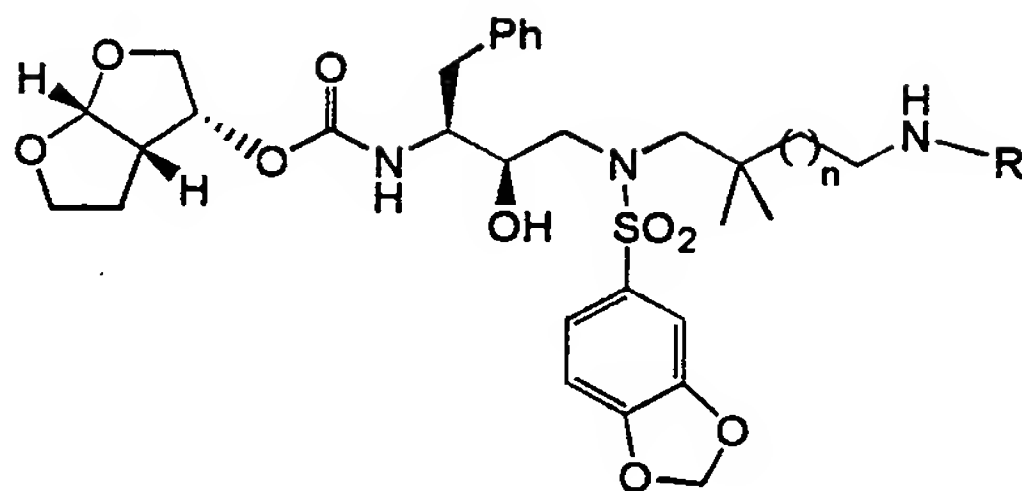
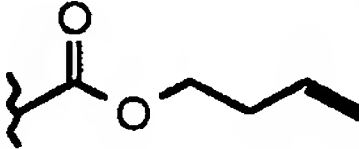
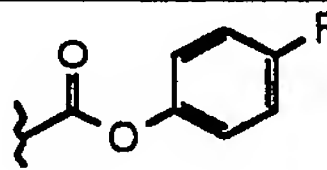
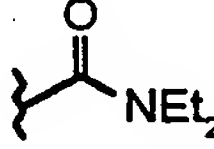
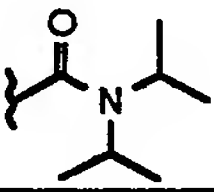
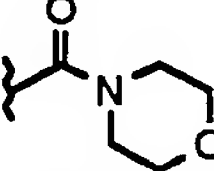
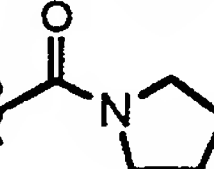
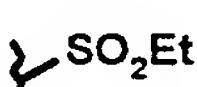
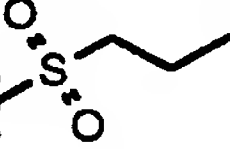
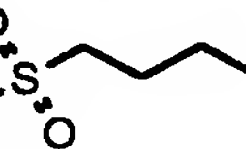
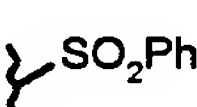
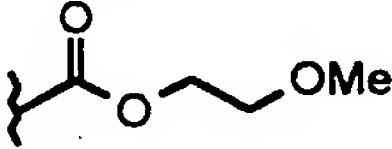
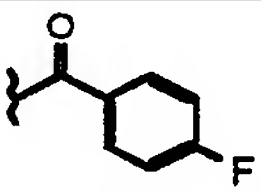
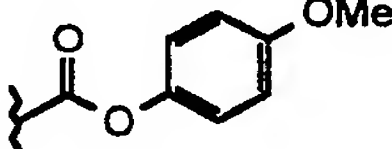
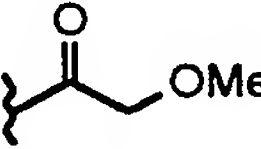
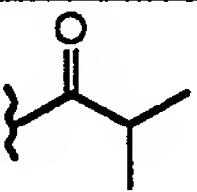
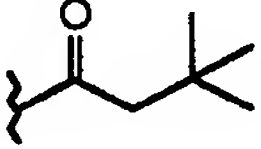
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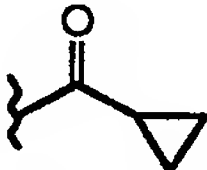
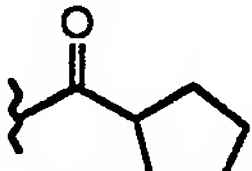
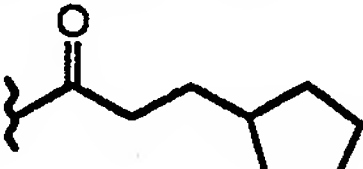
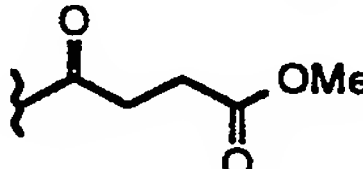
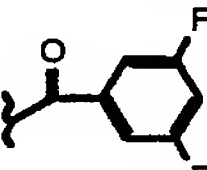
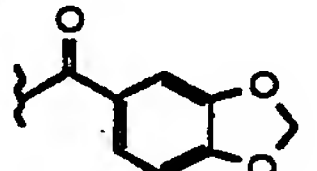
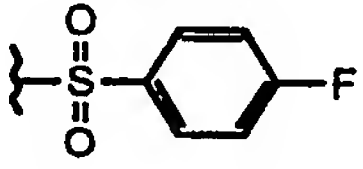
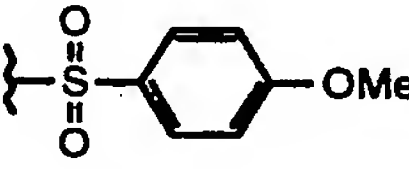
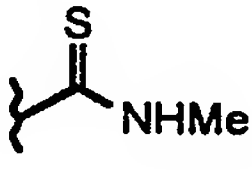
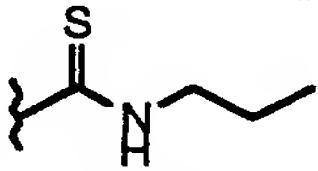
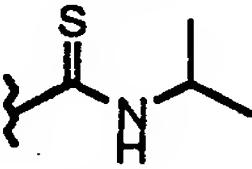
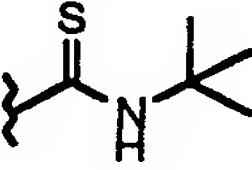
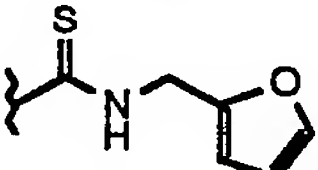
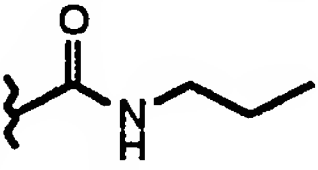
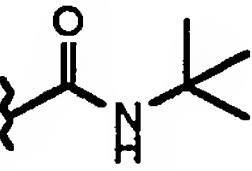
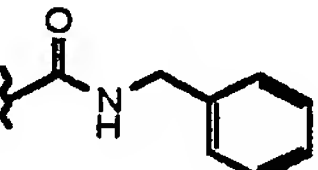
Table 5

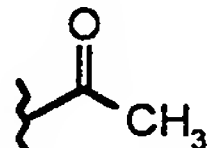
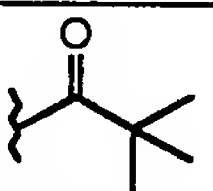
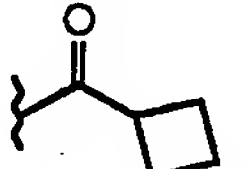
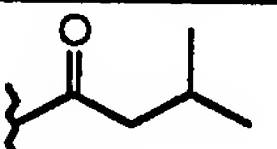
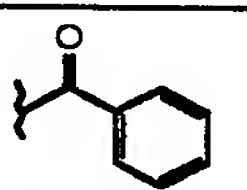
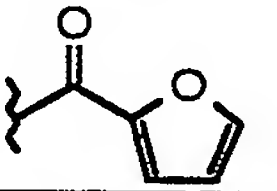
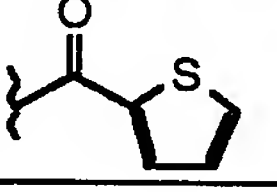
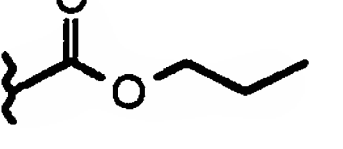
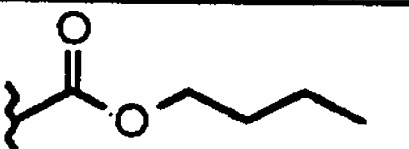
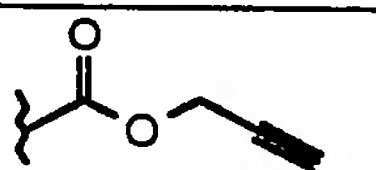
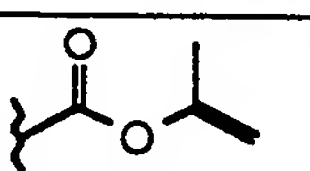
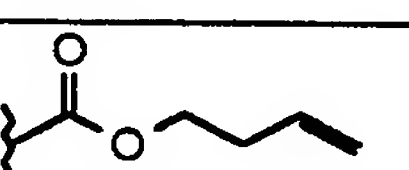
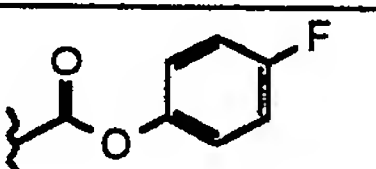
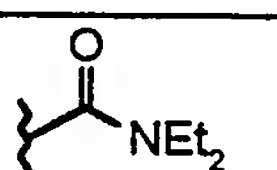
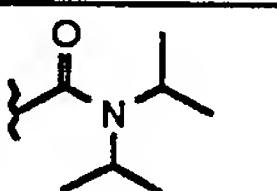
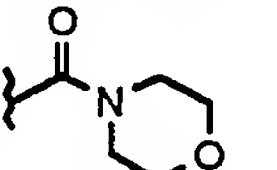
5

Compound	n	R
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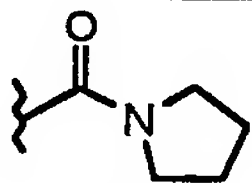
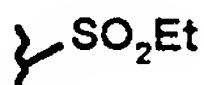
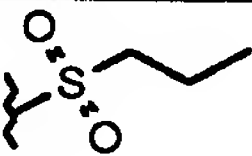
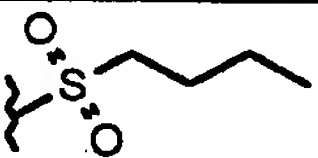
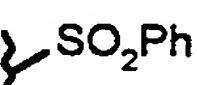
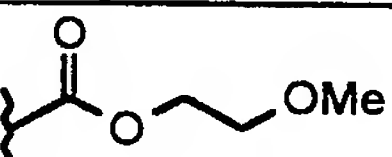
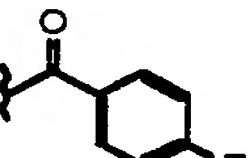
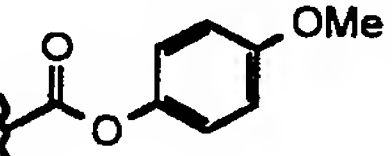
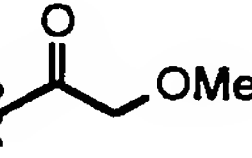
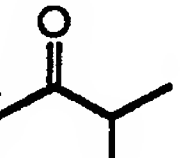
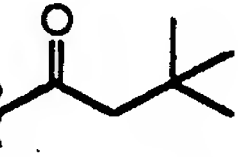
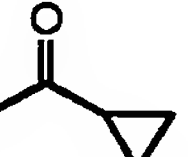
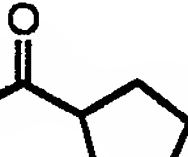
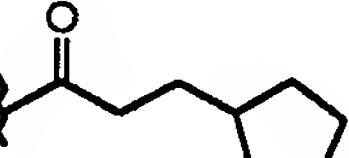
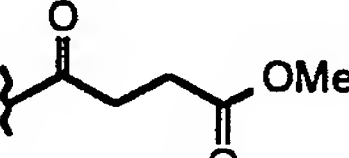
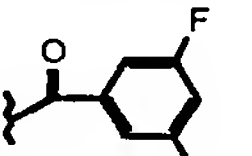
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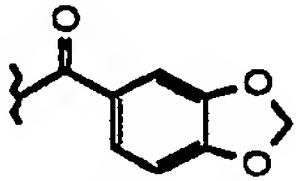
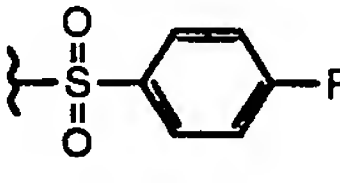
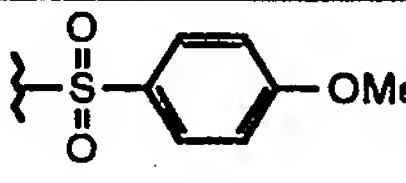
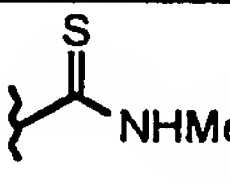
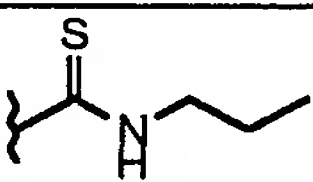
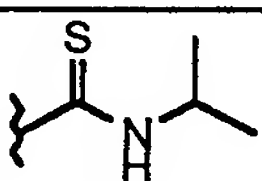
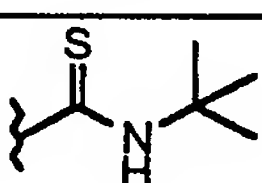
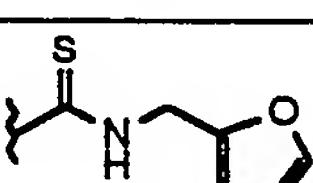
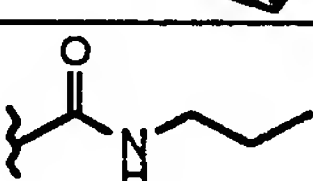
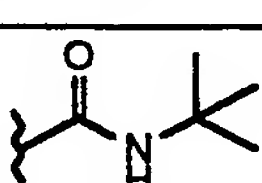
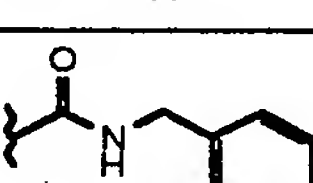
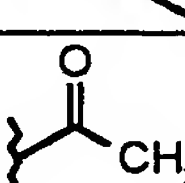
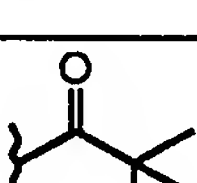
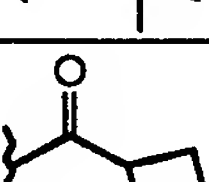
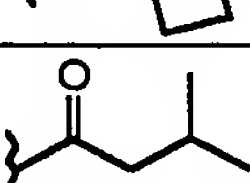
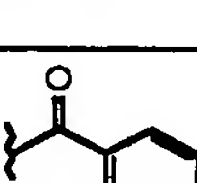
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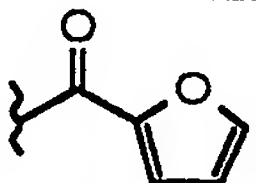
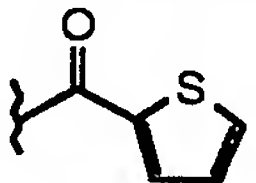
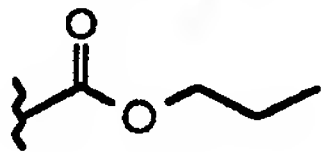
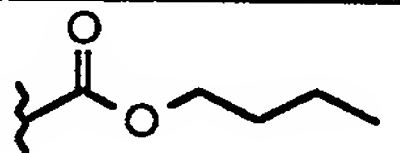
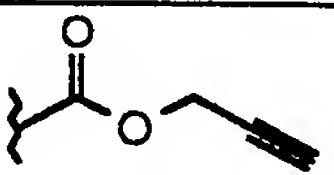
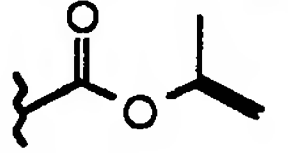
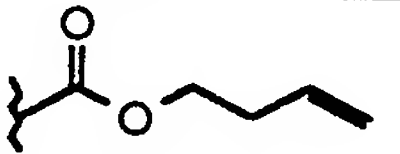
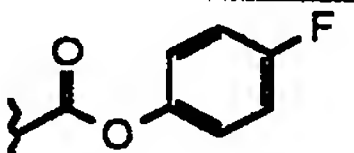
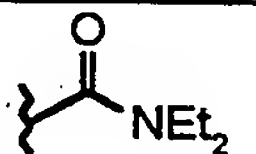
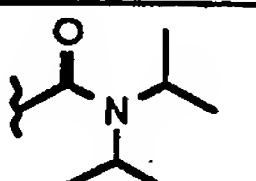
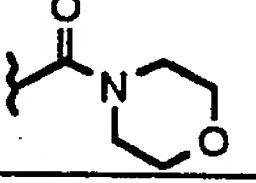
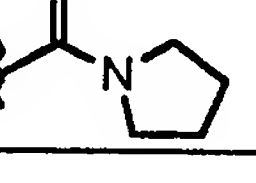
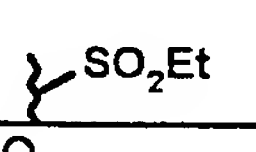


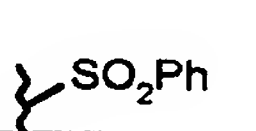
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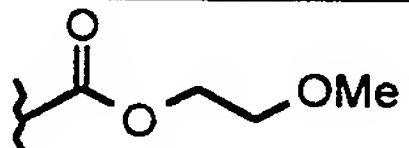
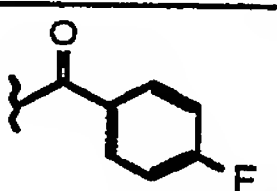
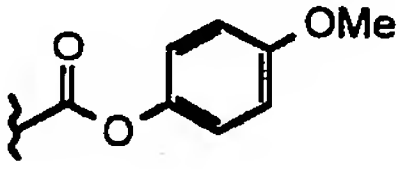
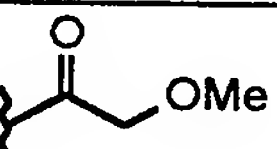
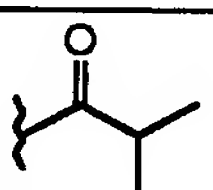
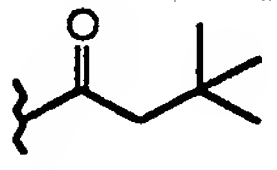
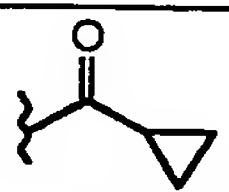
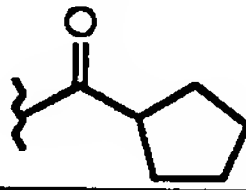
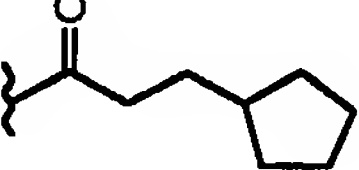
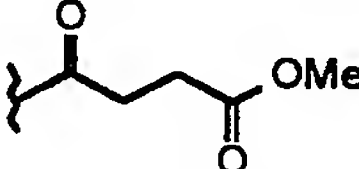
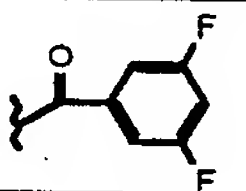
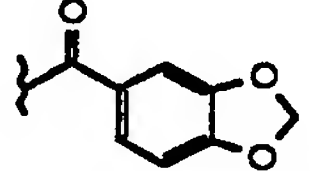
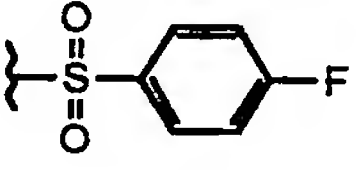
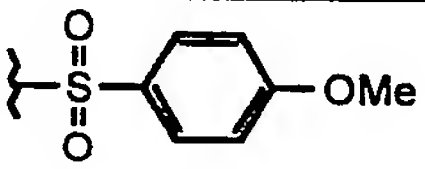
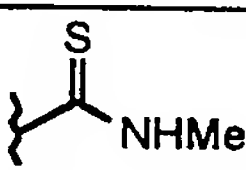
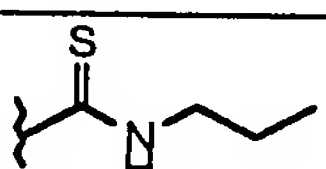
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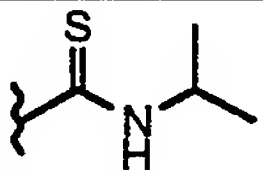
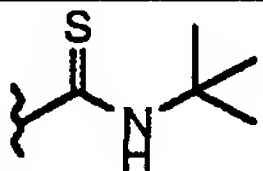
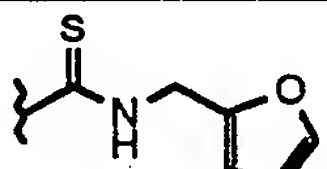
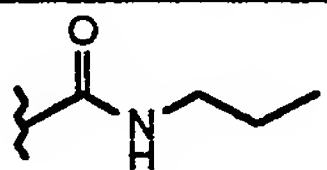
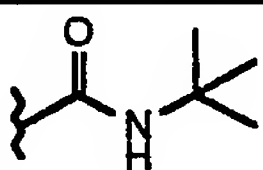
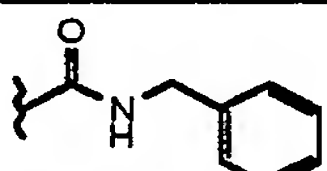
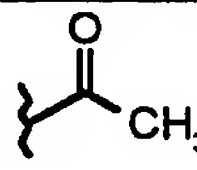
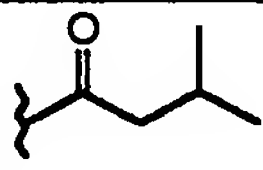
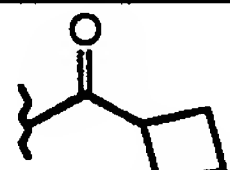
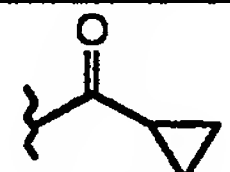
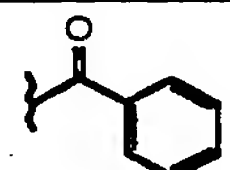
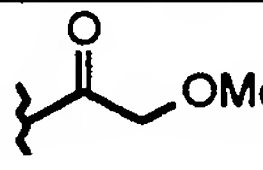
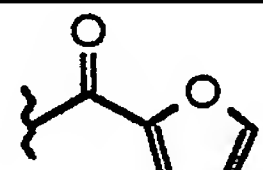
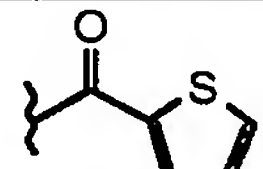
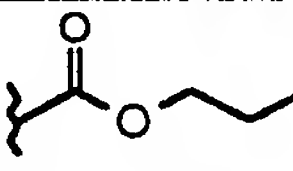
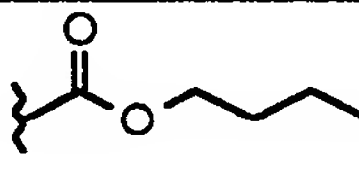
-56-

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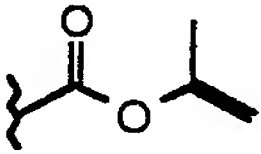
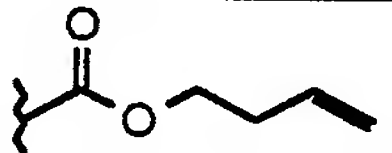
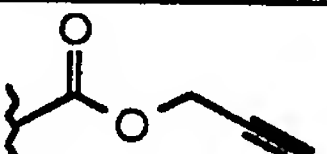
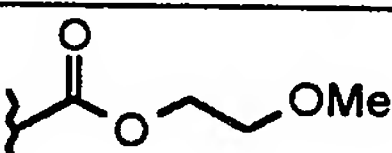
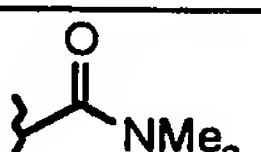
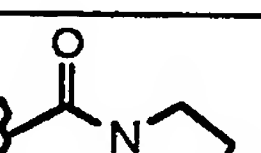
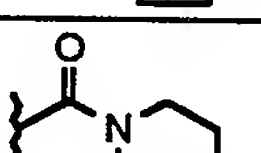
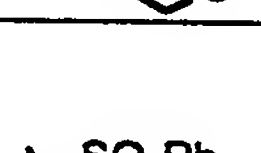
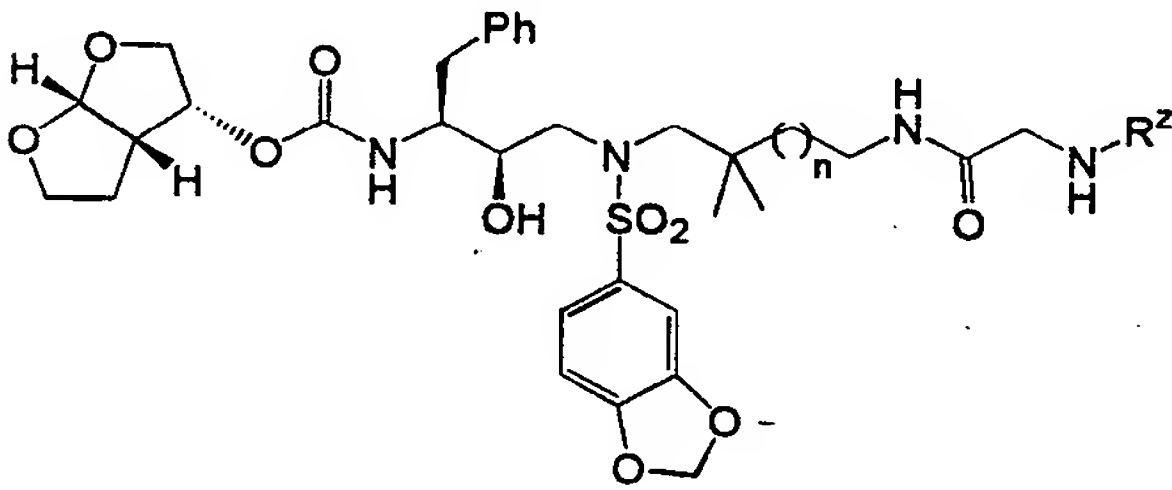
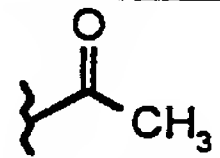
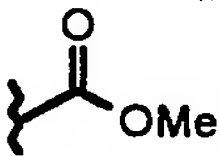
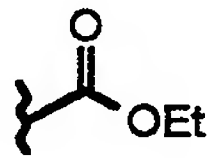
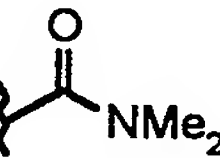
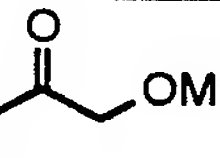

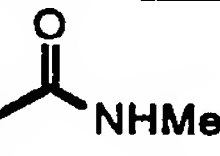
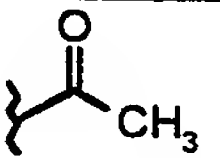
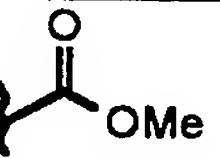
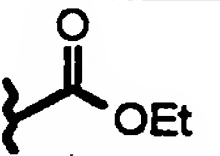
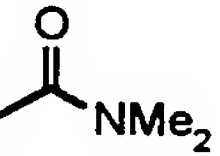
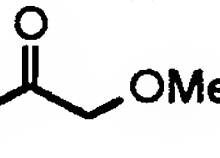


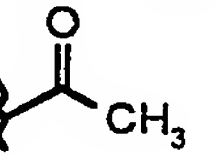
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519	4	

Table 6



5

Compound	n	R ^z
520	1	H
521	2	H

522	3	H
523	1	
524	1	
525	1	
526	1	
527	1	
528	1	
529	1	
530	2	
531	2	
532	2	
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535	2	
536	2	
537	3	

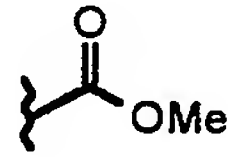
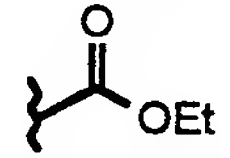
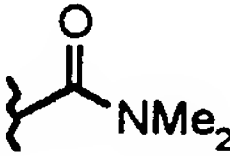
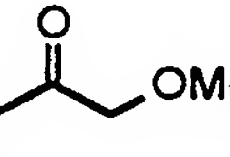
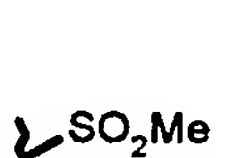
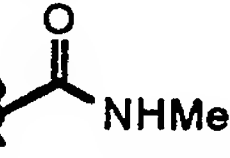
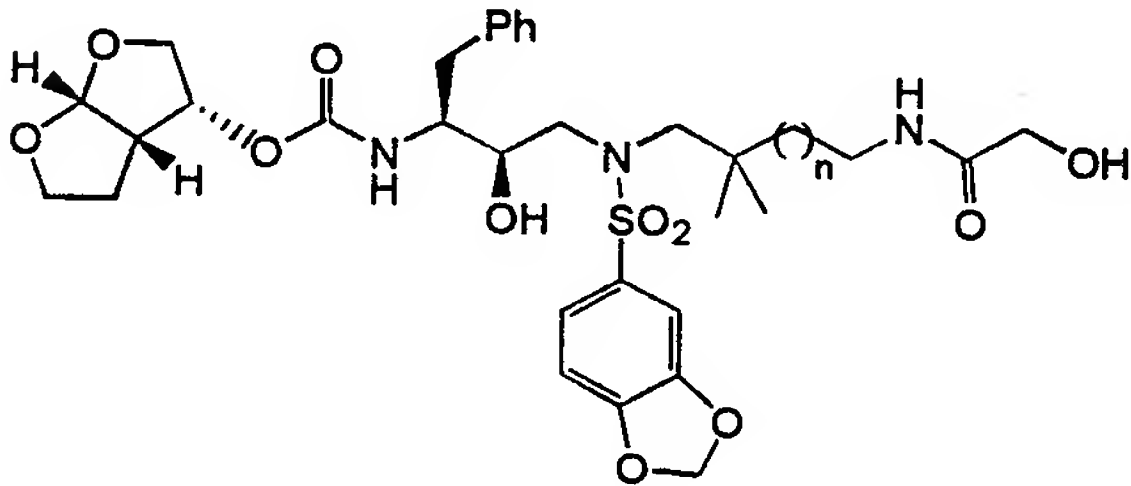
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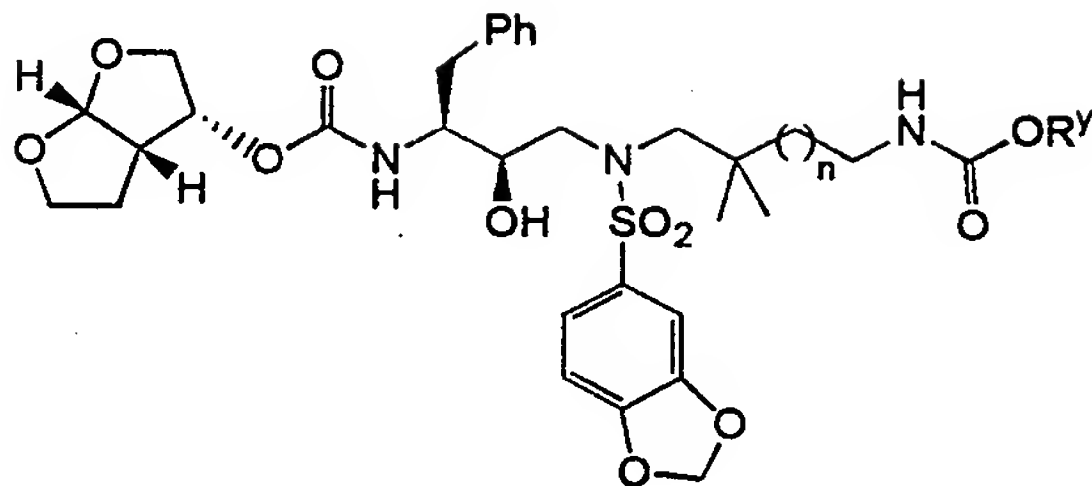
Table 7

5



Compound	N
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546	3
547	4

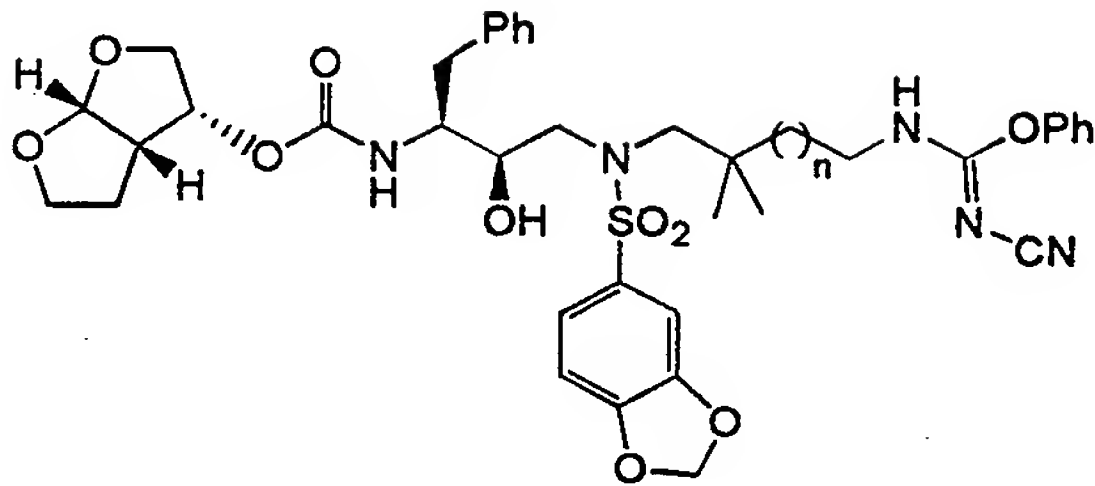
Table 8



5

Compound	n	R ^y
548	1	
549	1	
550	2	
551	2	
552	3	
553	3	
554	4	
555	4	

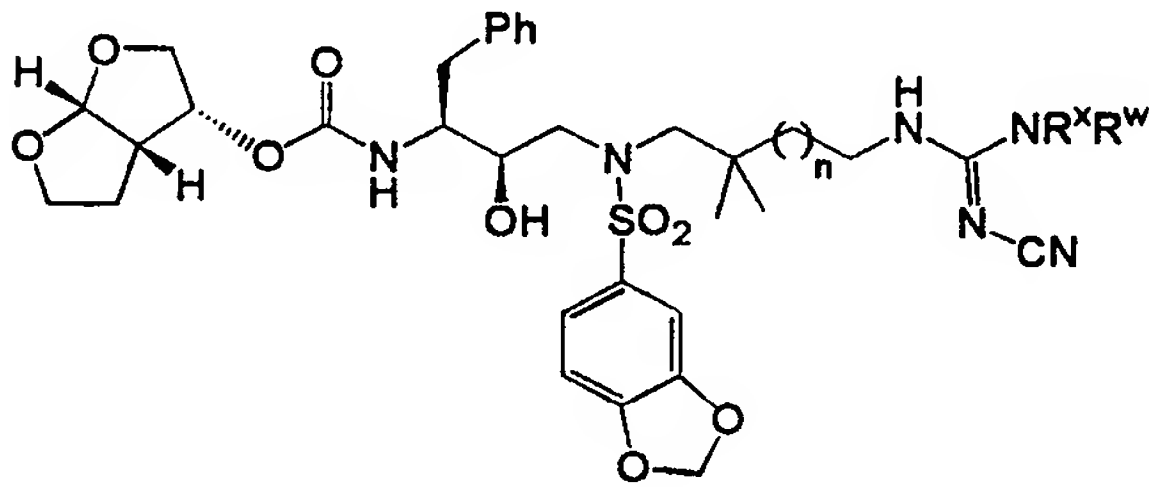
Table 9



5

Compound	n
556	1
557	2
558	3
559	4

Table 10

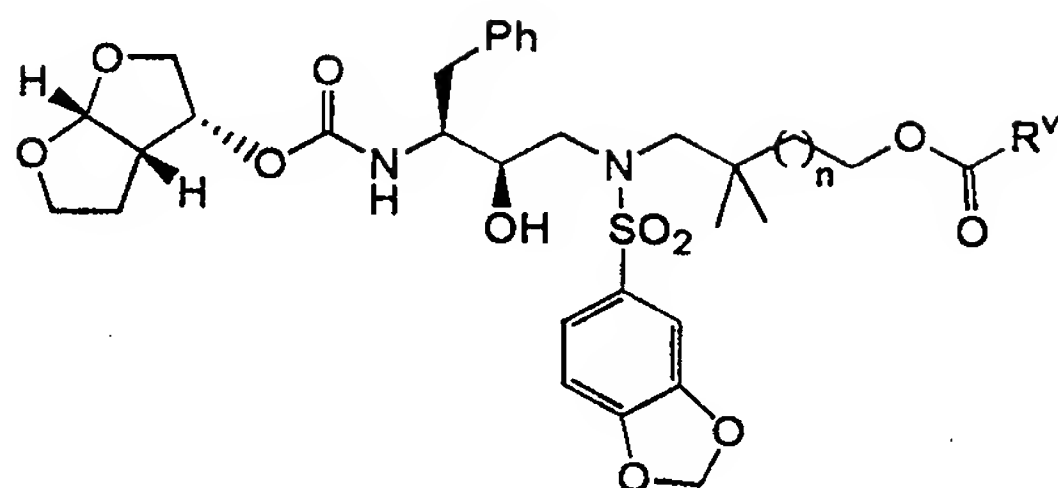


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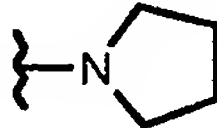
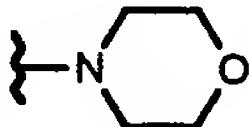



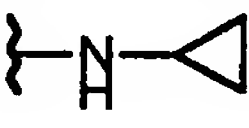
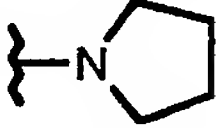



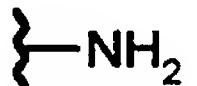



Compound	n	R ^x	R ^w
560	1	H	H
561	1	Me	H
562	1	n-Pr	H
563	1	Me	Me

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564	2	H	H
565	2	Me	H
566	2	n-Pr	H
567	2	Me	Me
568	3	H	H
569	3	Me	H
570	3	n-Pr	H
571	3	Me	Me
572	4	H	H
573	4	Me	H
574	4	n-Pr	H
575	4	Me	Me

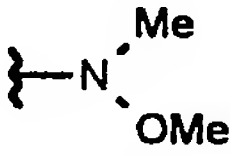
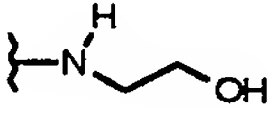
Table 11

Compound	n	R^v
576	1	
577	2	
578	3	
579	1	NMe_2
580	1	NH_2
581	1	NHEt
582	1	NHn-Pr
583	1	NHi-Pr
584	1	NHn-Bu
585	1	

586	1	
587	1	
588	1	
589	2	
590	2	
591	2	
592	2	
593	2	
594	2	
595	3	
596	3	
597	3	
598	3	
599	3	

600	3	
601	3	
602	3	
603	3	
604	3	
605	1	
606	1	
607	1	
608	1	
609	2	
610	2	
611	2	
612	2	
613	3	
614	3	

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615	3	
616	3	

Preferred compounds are compound numbers: 12, 16, 25, 29, 30, 31, 35, 39, 41, 42, 47, 100, 124, 210, 224, 240, 248, 250, 255, 263, 270, 272, 280, 299, 300, 307, 309, 313, 314, 315, 316, 359, 360, 375, 378, 384, 421, 459, 464, 483, 494, 496, 523, 524, 531, 542, 548, 553, 558, 563, 570, 571, 575, 579, 589, 596, 606, 609, 616.

In one embodiment of preferred compounds, the compounds are numbers: 210, 224, 240, 248, 250, 255, 263, 270, 272, 280, 299, 300, 307, 309, 313, 314, 315, 316, 359, 360, 384, 483, 494, 496, 523, 524, 531, 542, 548, 553, 558, 563, 570, 571, 575, 579, 589, 596, 606, 609, 616.

In another embodiment of preferred compounds, the compounds are numbers: 12, 16, 25, 29, 30, 31, 35, 39, 41, 42, 47, 100, 124, 375, 378, 421, 459, 464.

More preferred are compound numbers 12, 16, 25, 35, 39, 42, 47, 100, 224, 240, 263, 270, 272, 280, 299, 300, 307, 309, 313, 314, 315, 316, 359, 360, 375, 378, 384, 421, 459, 464, 483, 494, 496, 548, 553, 558, 563, 570, 571, 575, 579, 589, 596, 606, 609, 616.

In one embodiment of more preferred compounds, the compounds are numbers: 224, 240, 263, 270, 272, 280, 299, 300, 307, 309, 313, 314, 315, 316, 359, 360, 384, 483, 494, 496, 548, 553, 558, 563, 570, 571, 575, 579, 589, 596, 606, 609, 616.

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In another embodiment of more preferred compounds, the compounds are numbers: 12, 16, 25, 35, 39, 42, 47, 100, 375, 378, 421, 459, 464.

The most preferred compounds are compound
5 numbers 16, 25, 42, 47, 100, 224, 240, 272, 299, 314.

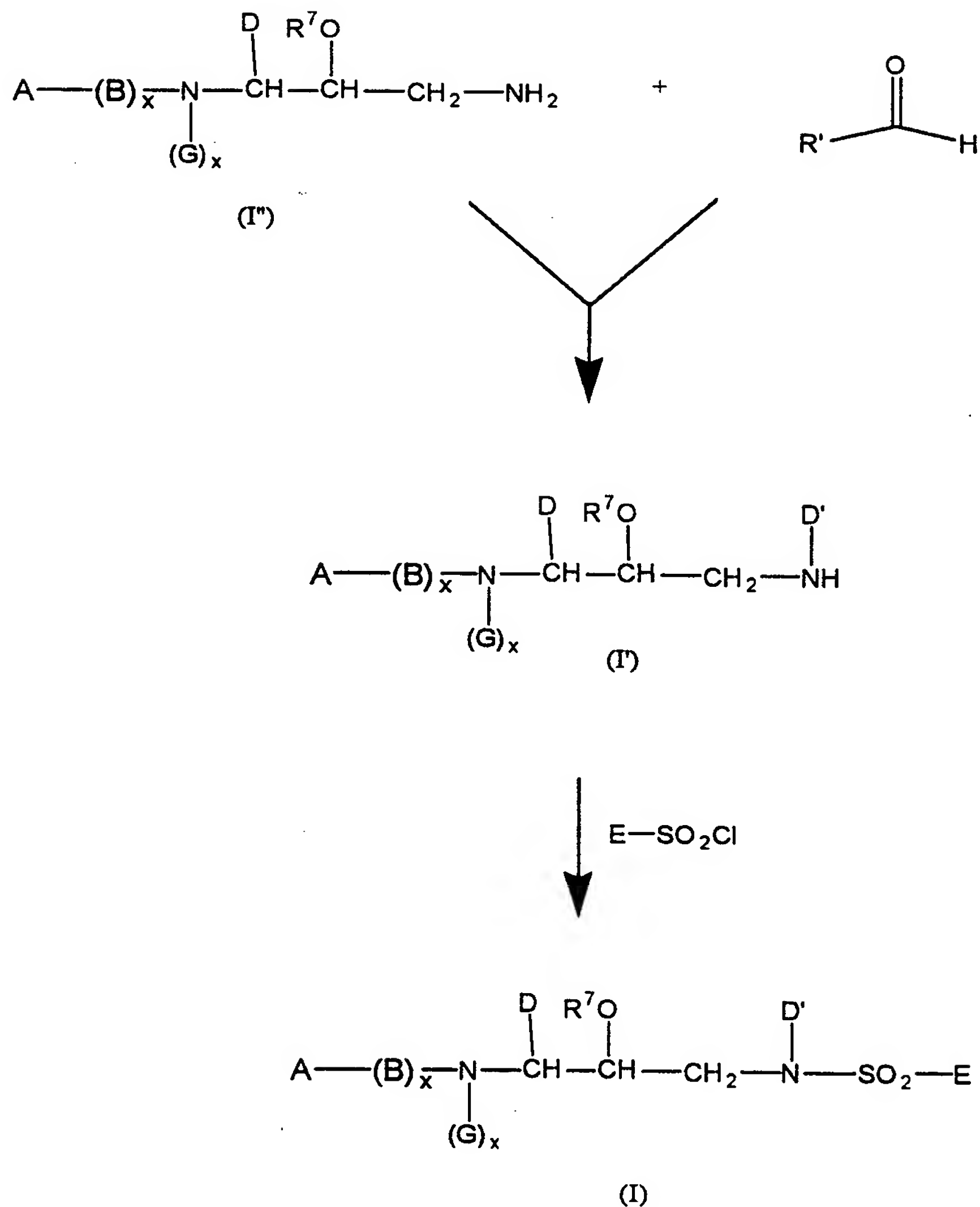
In one embodiment of most preferred compounds, the compounds are numbers: 224, 240, 272, 299, 314.

In another embodiment of most preferred compounds, the compounds are numbers: 16, 25, 42, 47,
10 100.

The compounds of the present invention can be readily prepared by techniques known in the art. Scheme I illustrates a general synthetic route to compounds of formula (I).

15

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SCHEME I

In Step 1 of Scheme I, radical R' is selected such that $R'-CH_2-$ is D' .

5

The synthetic approach illustrated in Scheme I can be readily extended to produce other compounds of the present invention. The above synthetic scheme is not intended to comprise a comprehensive list of all means by which compounds described and claimed in this application

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may be synthesized. Further methods will be evident to those of ordinary skill in the art.

As discussed above, the novel compounds of the present invention are excellent ligands for aspartyl proteases, particularly HIV-1 and HIV-2 proteases. Accordingly, these compounds are capable of targeting and inhibiting late stage events in HIV replication, i.e., the processing of the viral polyproteins by HIV encoded proteases. Such compounds inhibit the proteolytic processing of viral polyprotein precursors by inhibiting aspartyl protease. Because aspartyl protease is essential for the production of mature virions, inhibition of that processing effectively blocks the spread of virus by inhibiting the production of infectious virions, particularly from chronically infected cells. Compounds according to this invention advantageously inhibit the ability of the HIV-1 virus to infect immortalized human T cells over a period of days, as determined by an assay of extracellular p24 antigen -- a specific marker of viral replication. Other anti-viral assays have confirmed the potency of these compounds.

The compounds of this invention may be employed in a conventional manner for the treatment of viruses, such as HIV and HTLV, which depend on aspartyl proteases for obligatory events in their life cycle. Such methods of treatment, their dosage levels and requirements may be selected by those of ordinary skill in the art from available methods and techniques. For example, a compound of this invention may be combined with a pharmaceutically acceptable adjuvant for administration to a virally-infected patient in a

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pharmaceutically acceptable manner and in an amount effective to lessen the severity of the viral infection.

Alternatively, the compounds of this invention may be used in vaccines and methods for
5 protecting individuals against viral infection over an extended period of time. The compounds may be employed in such vaccines either alone or together with other compounds of this invention in a manner consistent with the conventional utilization of protease inhibitors in
10 vaccines. For example, a compound of this invention may be combined with pharmaceutically acceptable adjuvants conventionally employed in vaccines and administered in prophylactically effective amounts to protect individuals over an extended period time against HIV infection. As
15 such, the novel protease inhibitors of this invention can be administered as agents for treating or preventing HIV infection in a mammal.

The compounds of formula I, especially those having a molecular weight of less than about 700 g/mole,
20 may be readily absorbed by the bloodstream of mammals upon oral administration. Compounds of formula I having a molecular weight of less than about 600 g/mole are most likely to demonstrate oral availability. This surprisingly impressive oral availability makes such
25 compounds excellent agents for orally-administered treatment and prevention regimens against HIV infection.

The compounds of this invention may be administered to a healthy or HIV-infected patient either as a single agent or in combination with other anti-viral
30 agents which interfere with the replication cycle of HIV. By administering the compounds of this invention with other anti-viral agents which target different events in

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the viral life cycle, the therapeutic effect of these compounds is potentiated. For instance, the co-administered anti-viral agent can be one which targets early events in the life cycle of the virus, such as cell entry, reverse transcription and viral DNA integration into cellular DNA. Anti-HIV agents targeting such early life cycle events include, didanosine (ddI), alcitabine (ddC), d4T, zidovudine (AZT), polysulfated polysaccharides, sT4 (soluble CD4), ganiclovir, dideoxycytidine, trisodium phosphonoformate, eflornithine, ribavirin, acyclovir, alpha interferon and trimenotrexate. Additionally, non-nucleoside inhibitors of reverse transcriptase, such as TIBO or nevirapine, may be used to potentiate the effect of the compounds of this invention, as may viral uncoating inhibitors, inhibitors of trans-activating proteins such as tat or rev, or inhibitors of the viral integrase.

Combination therapies according to this invention exert a synergistic effect in inhibiting HIV replication because each component agent of the combination acts on a different site of HIV replication. The use of such combinations also advantageously reduces the dosage of a given conventional anti-retroviral agent which would be required for a desired therapeutic or prophylactic effect as compared to when that agent is administered as a monotherapy. These combinations may reduce or eliminate the side effects of conventional single anti-retroviral agent therapies while not interfering with the anti-retroviral activity of those agents. These combinations reduce potential of resistance to single agent therapies, while minimizing any associated toxicity. These combinations may also

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increase the efficacy of the conventional agent without increasing the associated toxicity. In particular, we have discovered that these compounds act synergistically in preventing the replication of HIV in human T cells.

5 Preferred combination therapies include the administration of a compound of this invention with AZT, ddI, ddC or d4T.

Alternatively, the compounds of this invention may also be co-administered with other HIV
10 protease inhibitors such as Ro 31-8959 (Roche), L-735,524 (Merck), XM 323 (Du-Pont Merck) and A-80,987 (Abbott) to increase the effect of therapy or prophylaxis against various viral mutants or members of other HIV quasi species.

15 We prefer administering the compounds of this invention as single agents or in combination with retroviral reverse transcriptase inhibitors, such as derivatives of AZT, or other HIV aspartyl protease inhibitors. We believe that the co-administration of the
20 compounds of this invention with retroviral reverse transcriptase inhibitors or HIV aspartyl protease inhibitors may exert a substantial synergistic effect, thereby preventing, substantially reducing, or completely eliminating viral infectivity and its associated
25 symptoms.

The compounds of this invention can also be administered in combination with immunomodulators (e.g., bropirimine, anti-human alpha interferon antibody, IL-2, GM-CSF, methionine enkephalin, interferon alpha,
30 diethyldithiocarbamate, tumor necrosis factor, naltrexone and rEPO); and antibiotics (e.g., pentamidine

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isethiorate) to prevent or combat infection and disease associated with HIV infections, such as AIDS and ARC.

When the compounds of this invention are administered in combination therapies with other agents, they may be administered sequentially or concurrently to the patient. Alternatively, pharmaceutical or prophylactic compositions according to this invention may be comprised of a combination of an aspartyl protease inhibitor of this invention and another therapeutic or prophylactic agent.

Although this invention focuses on the use of the compounds disclosed herein for preventing and treating HIV infection, the compounds of this invention can also be used as inhibitory agents for other viruses which depend on similar aspartyl proteases for obligatory events in their life cycle. These viruses include, as well as other AIDS-like diseases caused by retroviruses, such as simian immunodeficiency viruses, but are not limited to, HTLV-I and HTLV-II. In addition, the compounds of this invention may also be used to inhibit other aspartyl proteases, and in particular, other human aspartyl proteases, including renin and aspartyl proteases that process endothelin precursors.

Pharmaceutical compositions of this invention comprise any of the compounds of the present invention, and pharmaceutically acceptable salts thereof, with any pharmaceutically acceptable carrier, adjuvant or vehicle. Pharmaceutically acceptable carriers, adjuvants and vehicles that may be used in the pharmaceutical compositions of this invention include, but are not limited to, ion exchangers, alumina, aluminum stearate, lecithin, serum proteins, such as human serum albumin,

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buffer substances such as phosphates, glycine, sorbic acid, potassium sorbate, partial glyceride mixtures of saturated vegetable fatty acids, water, salts or electrolytes, such as protamine sulfate, disodium
5 hydrogen phosphate, potassium hydrogen phosphate, sodium chloride, zinc salts, colloidal silica, magnesium trisilicate, polyvinyl pyrrolidone, cellulose-based substances, polyethylene glycol, sodium carboxymethylcellulose, polyacrylates, waxes,
10 polyethylene-polyoxypropylene-block polymers, polyethylene glycol and wool fat.

The pharmaceutical compositions of this invention may be administered orally, parenterally, by inhalation spray, topically, rectally, nasally, buccally,
15 vaginally or via an implanted reservoir. We prefer oral administration or administration by injection. The pharmaceutical compositions of this invention may contain any conventional non-toxic pharmaceutically-acceptable carriers, adjuvants or vehicles. The term parenteral as
20 used herein includes subcutaneous, intracutaneous, intravenous, intramuscular, intra-articular, intrasynovial, intrasternal, intrathecal, intralesional and intracranial injection or infusion techniques.

The pharmaceutical compositions may be in the
25 form of a sterile injectable preparation, for example, as a sterile injectable aqueous or oleaginous suspension. This suspension may be formulated according to techniques known in the art using suitable dispersing or wetting agents (such as, for example, Tween 80) and suspending
30 agents. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example,

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as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are mannitol, water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose, any bland fixed oil may be employed including synthetic mono- or diglycerides. Fatty acids, such as oleic acid and its glyceride derivatives are useful in the preparation of injectables, as are natural pharmaceutically-acceptable oils, such as olive oil or castor oil, especially in their polyoxyethylated versions. These oil solutions or suspensions may also contain a long-chain alcohol diluent or dispersant such as Ph. Helv or a similar alcohol.

The pharmaceutical compositions of this invention may be orally administered in any orally acceptable dosage form including, but not limited to, capsules, tablets, and aqueous suspensions and solutions. In the case of tablets for oral use, carriers which are commonly used include lactose and corn starch. Lubricating agents, such as magnesium stearate, are also typically added. For oral administration in a capsule form, useful diluents include lactose and dried corn starch. When aqueous suspensions are administered orally, the active ingredient is combined with emulsifying and suspending agents. If desired, certain sweetening and/or flavoring and/or coloring agents may be added.

The pharmaceutical compositions of this invention may also be administered in the form of suppositories for rectal administration. These compositions can be prepared by mixing a compound of this

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invention with a suitable non-irritating excipient which is solid at room temperature but liquid at the rectal temperature and therefore will melt in the rectum to release the active components. Such materials include, but are not limited to, cocoa butter, beeswax and polyethylene glycols.

Topical administration of the pharmaceutical compositions of this invention is especially useful when the desired treatment involves areas or organs readily accessible by topical application. For application topically to the skin, the pharmaceutical composition should be formulated with a suitable ointment containing the active components suspended or dissolved in a carrier. Carriers for topical administration of the compounds of this invention include, but are not limited to, mineral oil, liquid petroleum, white petroleum, propylene glycol, polyoxyethylene polyoxypropylene compound, emulsifying wax and water. Alternatively, the pharmaceutical composition can be formulated with a suitable lotion or cream containing the active compound suspended or dissolved in a carrier. Suitable carriers include, but are not limited to, mineral oil, sorbitan monostearate, polysorbate 60, cetyl esters wax, cetearyl alcohol, 2-octyldodecanol, benzyl alcohol and water. The pharmaceutical compositions of this invention may also be topically applied to the lower intestinal tract by rectal suppository formulation or in a suitable enema formulation. Topically-transdermal patches are also included in this invention.

The pharmaceutical compositions of this invention may be administered by nasal aerosol or inhalation. Such compositions are prepared according to

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techniques well-known in the art of pharmaceutical formulation and may be prepared as solutions in saline, employing benzyl alcohol or other suitable preservatives, absorption promoters to enhance bioavailability, 5 fluorocarbons, and/or other solubilizing or dispersing agents known in the art.

Dosage levels of between about 0.01 and about 100 mg/kg body weight per day, preferably between about 0.5 and about 50 mg/kg body weight per day of the active 10 ingredient compound are useful in the prevention and treatment of viral infection, including HIV infection. Typically, the pharmaceutical compositions of this invention will be administered from about 1 to about 5 times per day or alternatively, as a continuous infusion. 15 Such administration can be used as a chronic or acute therapy. The amount of active ingredient that may be combined with the carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. A typical 20 preparation will contain from about 5% to about 95% active compound (w/w). Preferably, such preparations contain from about 20% to about 80% active compound.

Upon improvement of a patient's condition, a maintenance dose of a compound, composition or 25 combination of this invention may be administered, if necessary. Subsequently, the dosage or frequency of administration, or both, may be reduced, as a function of the symptoms, to a level at which the improved condition is retained when the symptoms have been alleviated to the 30 desired level, treatment should cease. Patients may, however, require intermittent treatment on a long-term basis upon any recurrence of disease symptoms.

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As the skilled artisan will appreciate, lower or higher doses than those recited above may be required. Specific dosage and treatment regimens for any particular patient will depend upon a variety of factors, including
5 the activity of the specific compound employed, the age, body weight, general health status, sex, diet, time of administration, rate of excretion, drug combination, the severity and course of the infection, the patient's disposition to the infection and the judgment of the
10 treating physician.

The compounds of this invention are also useful as commercial reagents which effectively bind to aspartyl proteases, particularly HIV aspartyl protease. As commercial reagents, the compounds of this invention,
15 and their derivatives, may be used to block proteolysis of a target peptide or may be derivatized to bind to a stable resin as a tethered substrate for affinity chromatography applications. These and other uses which characterize commercial aspartyl protease inhibitors will
20 be evident to those of ordinary skill in the art.

As used herein, the compounds according to the invention are defined to include pharmaceutically acceptable derivatives or prodrugs thereof. A
"pharmaceutically acceptable derivative" or
25 "pharmaceutically acceptable prodrug" means any pharmaceutically acceptable salt, ester, salt of an ester, or other derivative of a compound of this invention which, upon administration to a recipient, is capable of providing (directly or indirectly) a compound
30 of this invention or an active metabolite or residue thereof. Particularly favored derivatives and prodrugs are those that increase the bioavailability of the

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compounds of this invention when such compounds are administered to a mammal (e.g., by allowing an orally administered compound to be more readily absorbed into the blood) or which enhance delivery of the parent
5 compound to a biological compartment (e.g., the brain or lymphatic system) relative to the parent species.

The compounds according to the invention may be used in the form of salts derived from inorganic or organic acids. Included among such acid salts, for
10 example, are the following: acetate, adipate, alginate, aspartate, benzoate, benzenesulfonate, bisulfate, butyrate, citrate, camphorate, camphorsulfonate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, fumarate, flucoheptanoate,
15 glycerophosphate, hemisulfate, heptanoate, hexanoate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxyethanesulfonate, lactate, maleate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, oxalate, pamoate, pectinate, persulfate,
20 phenylpropionate, picrate, pivalate, propionate, succinate, tartrate, thiocyanate, tosylate and undecanoate. Other acids, such as oxalic, while not in themselves pharmaceutically acceptable, may be employed in the preparation of salts useful as intermediates in
25 obtaining the compounds of the invention and their pharmaceutically acceptable acid addition salts.

Salts derived from appropriate bases include alkali metal (e.g. sodium), alkaline earth metal (e.g., magnesium), ammonium and $^+NW_4$ (wherein W is C_{1-4} alkyl).
30 Physiologically acceptable salts of a hydrogen atom or an amino group include salts of organic carboxylic acids such as acetic, lactic, tartaric, malic, isethionic,

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lactobionic and succinic acids; organic sulfonic acids such as methanesulfonic, ethanesulfonic, benzenesulfonic and p-toluenesulfonic acids and inorganic acids such as hydrochloric, sulfuric, phosphoric and sulfamic acids.

- 5 Physiologically acceptable salts of a compound with a hydroxy group include the anion of said compound in combination with a suitable cation such as Na^+ , NH_4^+ , and NW_4^+ (wherein W is a C_{1-4} alkyl group).

Pharmaceutically acceptable salts include salts
10 of organic carboxylic acids such as ascorbic, acetic, citric, lactic, tartaric, malic, maleic, isothionic, lactobionic, p-aminobenzoic and succinic acids; organic sulphonic acids such as methanesulphonic, ethanesulphonic, benzenesulphonic and p-toluenesulphonic
15 acids and inorganic acids such as hydrochloric, sulphuric, phosphoric, sulphamic and pyrophosphoric acids.

For therapeutic use, salts of the compounds according to the invention will be pharmaceutically
20 acceptable. However, salts of acids and bases which are non-pharmaceutically acceptable may also find use, for example, in the preparation or purification of a pharmaceutically acceptable compound.

Preferred salts include salts formed from
25 hydrochloric, sulfuric, acetic, succinic, citric and ascorbic acids.

Preferred esters of the compounds according to the invention are independently selected from the following groups: (1) carboxylic acid esters obtained by
30 esterification of the hydroxy groups, in which the non-carbonyl moiety of the carboxylic acid portion of the ester grouping is selected from straight or branched

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chain alkyl (for example, acetyl, n-propyl, t-butyl, or n-butyl), alkoxyalkyl (for example, methoxymethyl), aralkyl (for example, benzyl), aryloxyalkyl (for example, phenoxymethyl), aryl (for example, phenyl optionally substituted by, for example, halogen, C₁₋₄alkyl, or C₁₋₄alkoxy or amino); (2) sulfonate esters, such as alkyl- or aralkylsulfonyl (for example, methanesulfonyl); (3) amino acid esters (for example, L-valyl or L-isoleucyl); (4) phosphonate esters and (5) mono-, di- or triphosphate esters. The phosphate esters may be further esterified by, for example, a C₁₋₂₀ alcohol or reactive derivative thereof, or by a 2,3-di (C₆₋₂₄)acyl glycerol.

In such esters, unless otherwise specified, any alkyl moiety present advantageously contains from 1 to 18 carbon atoms, particularly from 1 to 6 carbon atoms, more particularly from 1 to 4 carbon atoms. Any cycloalkyl moiety present in such esters advantageously contains from 3 to 6 carbon atoms. Any aryl moiety present in such esters advantageously comprises a phenyl group.

Any reference to any of the above compounds also includes a reference to a pharmaceutically acceptable salts thereof.

The compounds according to the invention are especially useful for the treatment of AIDS and related clinical conditions such as AIDS related complex (ARC), progressive generalized lymphadenopathy (PGL), Kaposi's sarcoma, thrombocytopenic purpura, AIDS-related neurological conditions such as AIDS dementia complex, multiple sclerosis or tropical paraperesis, and also anti-HIV antibody-positive and HIV-positive conditions, including such conditions in asymptomatic patients.

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In a further aspect of the invention there are provided the compounds according to the invention for use in medical therapy particularly for the treatment or prophylaxis of viral infections such as HIV infections.

5 According to another aspect, the present invention provides a method for the treatment or prevention of the symptoms or effects of a viral infection in an infected animal, for example, a mammal including a human, which comprises treating said animal
10 with a therapeutically effective amount of a compound according to the invention. According to a particular embodiment of this aspect of the invention, the viral infection is an HIV infection. A further aspect of the invention includes a method for the treatment or
15 prevention of the symptoms or effects of an HBV infection.

The compounds according to the invention may also be used in adjuvant therapy in the treatment of HIV infections or HIV-associated symptoms or effects, for
20 example Kaposi's sarcoma.

The present invention further provides a method for the treatment of a clinical condition in an animal, for example, a mammal including a human which clinical condition includes those which have been discussed in the
25 introduction hereinbefore, which comprises treating said animal with a therapeutically effective amount of a compound according to the invention. The present invention also includes a method for the treatment or prophylaxis of any of the aforementioned infections or
30 conditions.

In yet a further aspect, the present invention provides the use of a compound according to the invention

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in the manufacture of a medicament for the treatment or prophylaxis of any of the above mentioned viral infections or conditions. It will be appreciated that of compounds of Formula (I), (IA), (IB), (II), (III), (IV),
5 and (V) and one or more other HIV protease inhibitors, reverse transcriptase inhibitors, or non-nucleoside reverse transcriptase inhibitors may be used in the manufacture of the above medicament.

Reference herein to treatment extends to
10 prophylaxis as well as the treatment of established infections or symptoms.

The above compounds according to the invention and their pharmaceutically acceptable derivatives may be employed in combination with other therapeutic agents for
15 the treatment of the above infections or conditions. Combination therapies according to the present invention comprise the administration of at least one compound of the formula (I) or a pharmaceutically acceptable derivative thereof and at least one other
20 pharmaceutically active ingredient. The active ingredient(s) and pharmaceutically active agents may be administered simultaneously in either the same or different pharmaceutical formulations or sequentially in any order. The amounts of the active ingredient(s) and
25 pharmaceutically active agent(s) and the relative timings of administration will be selected in order to achieve the desired combined therapeutic effect. Preferably the combination therapy involves the administration of one compound according to the invention and one of the agents
30 mentioned herein below.

Examples of such further therapeutic agents include agents that are effective for the treatment of

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viral infections or associated conditions such as (1
alpha, 2 beta, 3 alpha)-9-[2,3-bis(hydroxymethyl)
cyclobutyl]guanine [(-)BHCG, SQ-34514], oxetanocin-G
(3,4-bis-(hydroxymethyl)-2-oxetanosyl]guanine), acyclic
5 nucleosides (e.g. acyclovir, valaciclovir, famciclovir,
ganciclovir, penciclovir), acyclic nucleoside
phosphonates (e.g. (S)-1-(3-hydroxy-2-phosphonyl-
methoxypropyl)cytosine (HPMPC), ribonucleotide reductase
inhibitors such as 2-acetylpyridine 5-[(2-
10 chloroanilino)thiocarbonyl) thiocarbonohydrazone,
3'-azido-3'-deoxythymidine, other 2',3'-dideoxynucleosides
such as 2',3'-dideoxycytidine, 2',3'-dideoxyadenosine,
2',3'-dideoxyinosine, 2',3'-didehydrothymidine, protease
inhibitors such as indinavir, ritonavir, nelfinavir, [3S-
15 [3R*(1R*, 2S*)]]-[3[[(4-aminophenyl)sulfonyl] (2-
methylpropyl)amino]-2-hydroxy-1-(phenylmethyl)propyl]-
tetrahydro-3-furanyl ester (141W94), oxathiolane
nucleoside analogues such as (-)-cis-1-(2-hydroxymethyl)-
1,3-oxathiolane 5-yl)-cytosine (lamivudine) or cis-1-(2-
20 (hydroxymethyl)-1,3-oxathiolan-5-yl)-5-fluorocytosine
(FTC), 3'-deoxy-3'-fluorothymidine, 5-chloro-2',3'-
dideoxy-3'-fluorouridine, (-)-cis-4-[2-amino-6-
(cyclopropylamino)-9H-purin-9-yl]-2-cyclopentene-1-
methanol, ribavirin, 9-[4-hydroxy-2-(hydroxymethyl)but-1-
25 yl]-guanine (H2G), tat inhibitors such as 7-chloro-5-(2-
pyrryl)-3H-1,4-benzodiazepin-2-(H)one (Ro5-3335), 7-
chloro-1,3-dihydro-5-(1H-pyrrol-2yl)-3H-1,4-
benzodiazepin-2-amine (Ro24-7429), interferons such as α -
interferon, renal excretion inhibitors such as
30 probenecid, nucleoside transport inhibitors such as
dipyridamole; pentoxifylline, N-acetylcysteine (NAC),
Procysteine, α -trichosanthin, phosphonoformic acid, as

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well as immunomodulators such as interleukin II or thymosin, granulocyte macrophage colony stimulating factors, erythropoietin, soluble CD₄ and genetically engineered derivatives thereof, or non-nucleoside reverse transcriptase inhibitors (NNRTIs) such as nevirapine (BI-5 5 RG-587), zalcitabine (α -APA) and delaviridine (BHAP), and phosphonoformic acid, and 1,4-dihydro-2H-3,1-benzoxazin-2-ones NNRTIs such as (-)-6-chloro-4-cyclopropylethynyl-4-trifluoromethyl-1,4-dihydro-2H-3,1-benzoxazin-2-one (L-10 743,726 or DMP-266), and quinoxaline NNRTIs such as isopropyl (2S)-7-fluoro-3,4-dihydro-2-ethyl-3-oxo-1(2H)-quinoxalinecarboxylate (HBY1293).

More preferably the combination therapy involves the administration of one of the above mentioned agents and a compound within one of the preferred or particularly preferred sub-groups within formula (I) as described above. Most preferably the combination therapy involves the joint use of one of the above named agents together with one of the compounds of formula (I) 20 specifically named herein.

The present invention further includes the use of a compound according to the invention in the manufacture of a medicament for simultaneous or sequential administration with at least one other therapeutic agent, such as those defined hereinbefore. 25

In order that this invention may be more fully understood, the following examples are set forth. These examples are for the purpose of illustration only and are not to be construed as limiting the scope of the invention in any way. 30

-90-

Example (Compound 1)

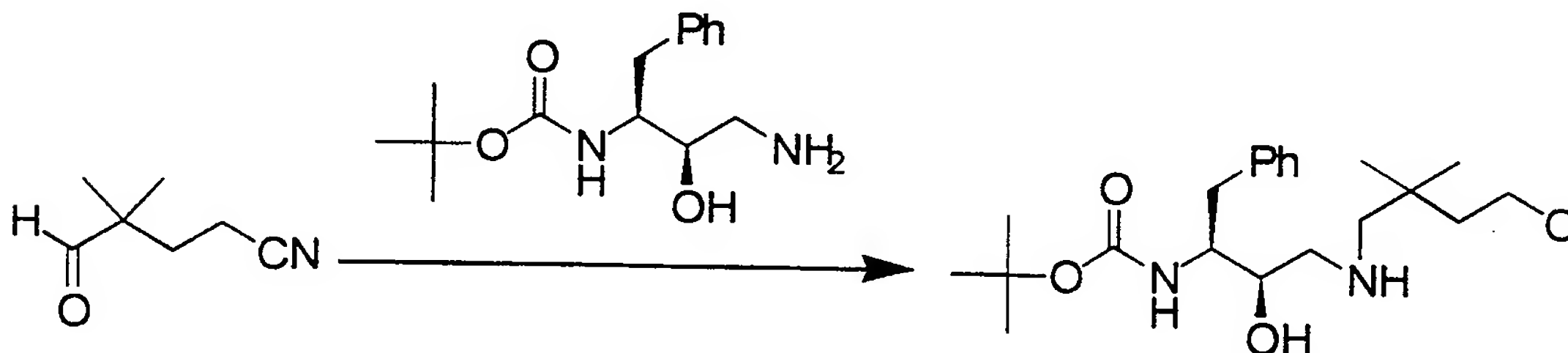
Step 1:

**3,3-Dimethyl-4-oxobutyl cyanide**

To a stirred solution of isobutyraldehyde (6.3 mL, 69.3 mmol), hydroquinone (25 mg) and acrylonitrile (5.7 mL, 86.7 mmol) in 1,4-dioxane (25 mL) was added 5% sodium hydroxide (4.6 mL). The reaction was heated at 65°C for 2.5 hours and then stirred at ambient temperature for 16 hours. The dioxane was removed *in vacuo* and the residue was taken up in dichloromethane (80 mL). The dichloromethane was washed with water (3 x 80 mL), dried (magnesium sulfate), concentrated *in vacuo* and then distilled to afford the title compound (5.4 g, 63%) as a colorless liquid; bp 70°C (0.8 mm);

¹H NMR (CDCl₃) : δ 1.11 (6H, s), 1.87 (2H, t), 2.29 (2H, t), 9.40 (1H, s)

Step 2:



tert-Butyl N-(1S,2R)-1-benzyl-3-[(4-cyano-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

To a suspension of the product of Step 1 (1.1 g, 8.7 mmol) and tert-butyl N-(1S,2R)-3-amino-1-benzyl-2-

-91-

hydroxypropylcarbamate (3.1 g, 11.0 mmol) in 1,2-dichloroethane (30 mL), tetrahydrofuran (30 mL) and *N,N*-dimethylformamide (30 mL) was added sodium triacetoxy borohydride (3.2 g, 15.0 mmol) and the mixture was

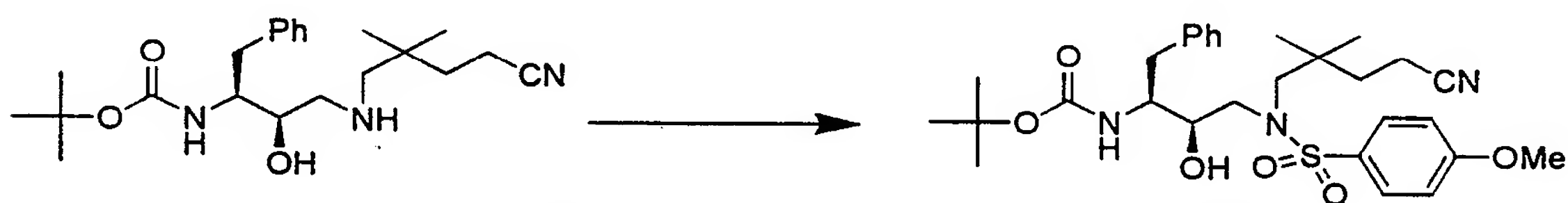
5 stirred under nitrogen atmosphere for 3.5 hours. The mixture was filtered and the filtrate was concentrated to 25 mL. The residue was partitioned between ethyl acetate and 2M sodium hydroxide/water. The organic phase was

10 washed with water, brine, dried (sodium sulfate), and evaporated. The residue was chromatographed (silica gel, dichloromethane/methanol, 95:5) to provide the title compound as a white solid (2.8 g, 90%); mp 79-80 C;

¹H NMR (DMSO-*d*₆): δ 0.81 (6H, s), 1.24 (9H, s), 1.50 (3H, br dd), 2.22 (2H, s), 2.40 (2H, br dd), 2.45-2.60 (3H, m), 2.96 (1H, dd), 3.40 (1H, br s), 3.50-3.55 (1H, m),

15 4.87 (1H, br s), 6.70 (1H, d), 7.10-7.18 (3H, m), 7.20-7.25 (2H, m); MS: 390.2 (MH⁺)

Step 3:



***tert*-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-(4-cyano-2,2-**

20 **dimethylbutyl) [(4-methoxyphenyl) sulfonyl]amino-2-**

hydroxypropyl) carbamate

To a solution of the product of Step 2 (0.5 g, 1.3 mmol) in anhydrous dichloromethane (15 mL) at 0°C was added *N*-ethyldiisopropylamine (0.48 mL, 0.36 g, 2.8 mmol) and a solution of 4-methoxybenzenesulfonyl chloride (0.27

25 g, 1.3 mmol) in anhydrous dichloromethane (2 mL) and the mixture was stirred at ambient temperature under nitrogen atmosphere for 18 hours. The mixture was washed

-92-

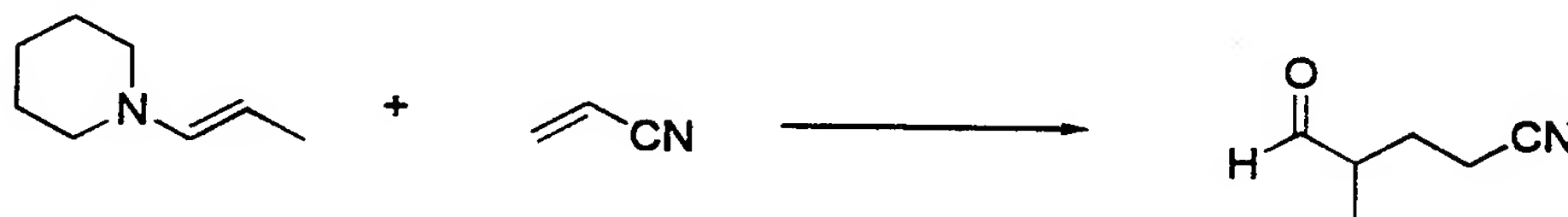
sequentially with 1M hydrochloric acid, sodium bicarbonate/water, brine, dried (sodium sulfate) and concentrated. The residue was chromatographed (silica gel, dichloromethane/methanol, 97:3) to provide the title compound as a colorless foam (0.34 g, 48%);

¹H NMR (DMSO-d₆): δ 0.88 (3H, s), 0.92 (3H, s), 1.20 (9H, s), 1.67 (2H, t), 2.40-2.50 (2H, m), 2.70 (1H, d), 2.80-2.95 (2H, m), 3.12-3.2 (4H, m), 3.73 (1H, br s), 3.82 (3H, s), 5.10 (1H, d), 6.63 (1H, d), 7.08 (2H, d), 7.10-7.27 (5H, m), 7.78 (2H, d); MS:560.2 (MH⁺)

Example (Compound 2)

Step 1:

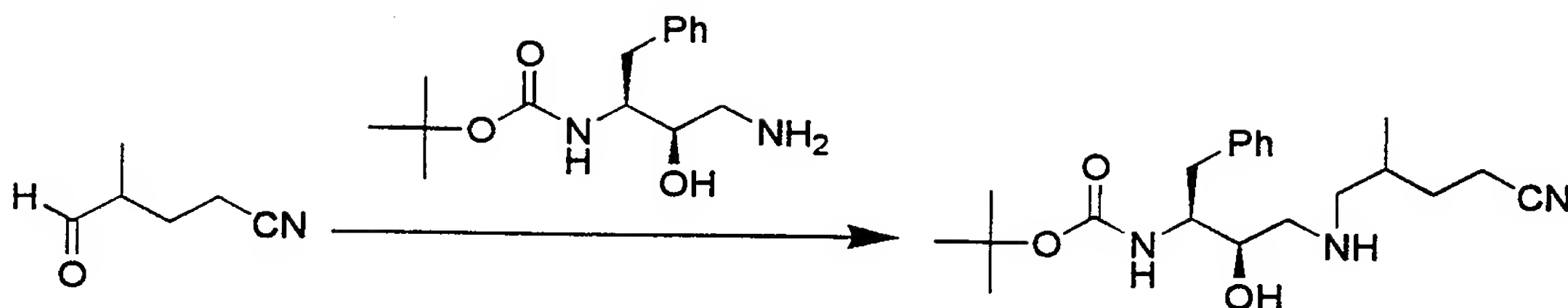
3-Methyl-4-oxobutyl cyanide



N-(1-propen-1-yl)piperidine [Mannich, D.; Ber. 1936, 69, 2106] was subjected to the procedure used in Step 1 of Example (Compound 3) to give the title compound as a pale yellow oil; bp 127-129 C (19 mm); ¹H NMR (CDCl₃): δ 1.19 (3H, d), 1.62-1.72 (1H, m), 2.03-2.13 (1H, m), 2.43 (2H, t), 2.51-2.61 (1H, m), 9.62 (1H, s).

Step 2:

-93-

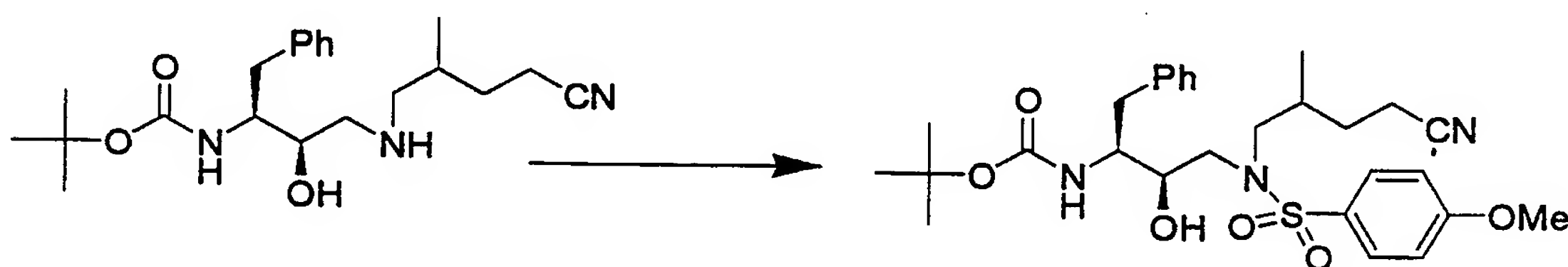


tert-Butyl *N*-(1*S*,2*R*)-1-benzyl-3-[(2*RS*)-(4-cyano-2-methylbutyl)amino]-2-hydroxypropylcarbamate

The product from step 1 was subjected to the procedure used in Example (Compound 1), Step 2, to provide the title compound as a white solid;

¹H NMR (DMSO-*d*₆): δ 0.93 (3H, d), 1.32 (9H, s), 1.34-1.84 (4H, m), 2.36-2.68 (7H, m), 3.04 (1H, dd), 3.42-3.66 (2H, m), 4.82 (1H, br s), 6.75 (1H, d), 7.15-7.35 (5H, m); MS: 376 (MH⁺)

Step 3:



ter-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-[(2*RS*)-(4-cyano-2-methylbutyl)][(4-methoxyphenyl)sulfonyl]amino)-2-hydroxypropylcarbamate

The product from Step 2 was subjected to the procedure used above (Step 3, Example (Compound 1)) to give the title compound as a white foam (~1:1 mixture of diastereomers by ¹H NMR spectroscopy);

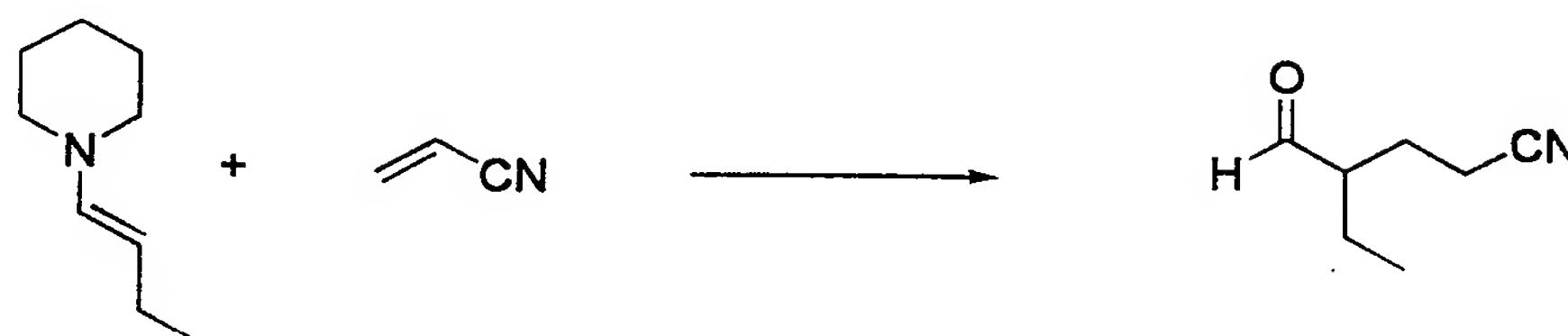
¹H NMR (DMSO-*d*₆): δ 0.80 (3H, d), 1.21 (9H, s), 1.2-1.3 (1H, m), 1.8-1.9 (2H, m), 2.35-2.45 (3H, m), 2.60-2.65 (1H, m), 2.70-2.82 (1H, m), 2.9-3.0 (2H, m), 3.22-3.30 (1H, m), 3.4-3.6 (2H, m), 3.8 (3H, m), 5.05 (1H, br s),

-94-

6.70 (1H, d), 7.05 (2H, d), 7.10-7.23 (5H, m), 7.70 (2H, d); MS: 546.3 (MH⁺)

Example (Compound 3)

Step 1:



5

3-Ethyl-4-oxobutyl cyanide

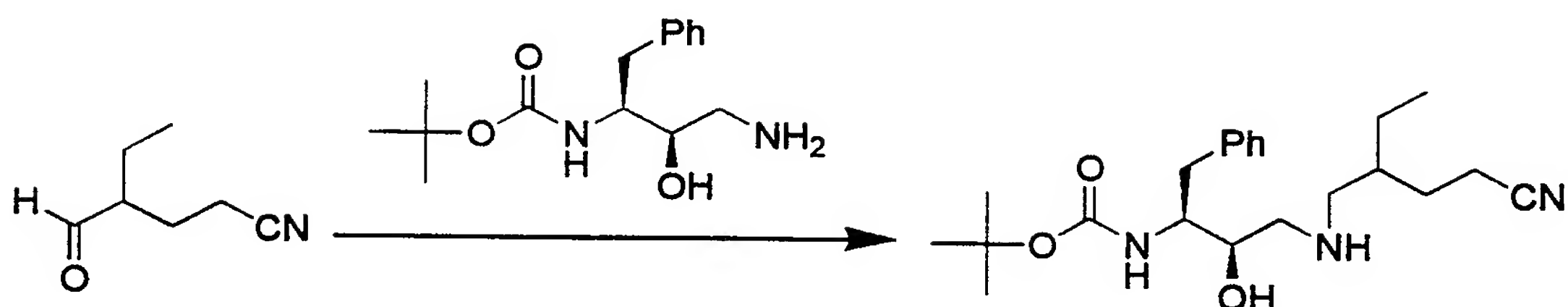
A stirred solution of N-(1-buten-1-yl)piperidine [Mannich, D.; Ber. 1936, 69, 2106] (4.2 g, 29.9 mmol) and acrylonitrile (2.1 g, 38.8 mmol) in acetonitrile (25 mL) was heated at reflux for 18 hours. Acetic acid and water (10 mL, 1:1) was added and the reaction was heated an additional 4 hours. The acetonitrile was removed *in vacuo* and the residue was taken up in ether (60 mL). The ether was washed with water (2 x 40 mL), saturated sodium bicarbonate/water (40 mL), dried (magnesium sulfate), concentrated *in vacuo* and then distilled to afford the title compound (1.6 g, 42%) as a colorless liquid. bp 132°C (18 mm);

¹H NMR (CDCl₃): δ 0.97 (3H, t), 1.55-1.82 (3H, m), 1.96-2.07 (1H, m), 2.40 (2H, q), 2.40-2.48 (1H, m), 9.62 (1H, s).

20

Step 2:

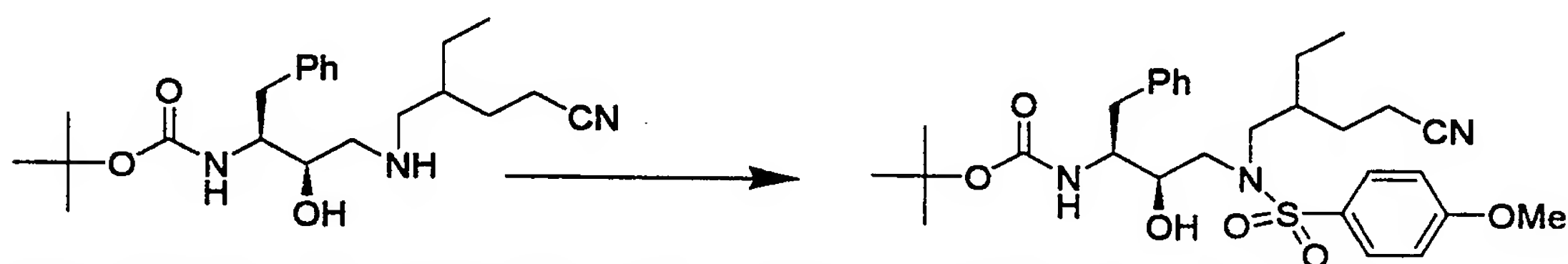
-95-



***tert*-Butyl *N*-(1*S*,2*R*)-1-benzyl-3-[(2*RS*)-(4-cyano-2-ethylbutyl)amino]-2-hydroxypropylcarbamate**

The product from step a was subjected to the general procedure (Step 2, Example (Compound 1)) to provide the title compound as a white solid; mp 67-68°C; ¹H NMR (DMSO-*d*₆): δ 0.90 (3H, t), 1.22 (9H, s), 1.23-1.28 (1H, m), 1.35-1.42 (1H, m), 1.45-1.62 (3H, m), 2.35-2.60 (7H, m), 2.96 (1H, dd), 3.40 (1H, br s), 3.5 (1H, qd), 4.80 (1H, br s), 6.73 (1H, d), 7.10-7.25 (5H, m); MS: 390.2 (MH⁺).

Step 3:



***tert*-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-[(2*RS*)-(4-cyano-2-ethylbutyl)][(4-methoxyphenyl)sulfonyl]amino)-2-hydroxypropyl)carbamate**

The product from step 2 was subjected to general procedure (Step 3, Example (Compound 1)) to give the title compound as a white glass (~1:1 mixture of diastereomers by ¹H NMR spectroscopy);

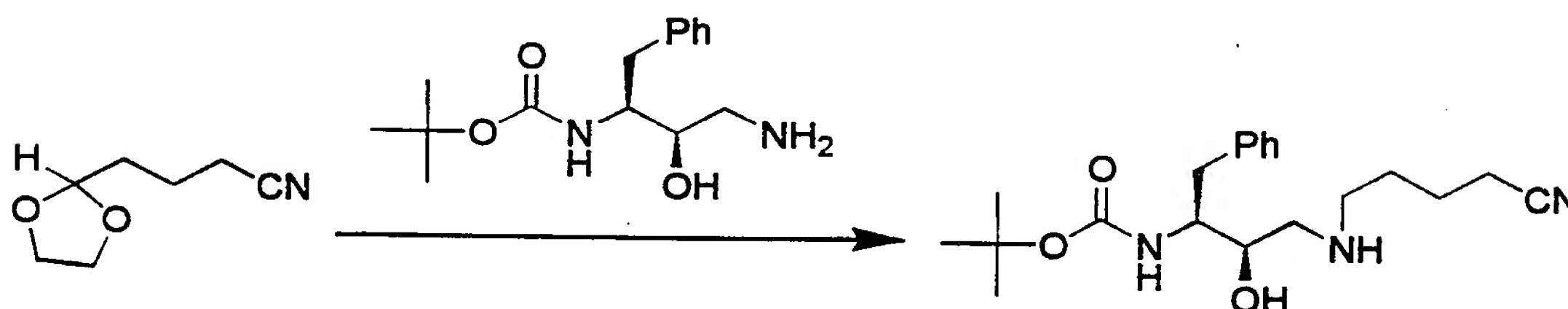
¹H NMR (DMSO-*d*₆): δ 0.80 (3H, t), 1.22 (9H, s), 1.32-1.40 (1H, m), 1.45-1.60 (1H, m), 1.75-1.82 (1H, m), 2.40-2.50

-96-

(3H, m), 2.75-2.85 (2H, m), 2.90 (1H, dd), 3.20 (1H, dd), 3.25-3.30 (1H, m), 3.40-3.48 (1H, m), 3.50-3.58 (1H, m), 3.80 (3H, s), 5.0 (1H, br s), 6.70 (1H, d), 7.08 (2H, d), 7.10-7.24 (5H, m), 7.75 (2H, m); MS: 582.3 (MH⁺)

5 Example (Compound 4)

Step 1:



3-(1,3-Dioxolan-2-yl)propionitrile

A mixture of 2-(3-chloropropyl)-1,3-dioxolane (5.7 g, 37.8 mmol) and potassium cyanide (3.8 g, 58.4 mmol) in dimethylsulfoxide (12 mL) was heated at 100 °C for 48 hours. The reaction mixture was diluted with water (80 mL) and extracted with ether (3 x 60 mL). The combined ether layer was washed with water (3 x 60 mL), dried (magnesium sulfate), concentrated in vacuo and purified by silica gel chromatography (ethyl acetate in hexane, 1:2) to afford the title compound (2.1g, 40%) as a pale yellow liquid;

¹H NMR (CDCl₃): δ 1.76-1.84 (4H, m), 4.21 (2H, t), 3.78-4.01 (4H, m), 4.87 (1H, t).

Step 2:

tert-Butyl *N*-(1*S*,2*R*)-1-benzyl-3-[(4-cyanobutyl)amino]-2-hydroxypropylcarbamate

-97-

To a stirred solution of the product from step 1 (2.0 g, 14.2 mmol) in tetrahydrofuran (85 mL) was added 5% hydrochloric acid (17 mL). The reaction was heated at 70 °C for 5 hours and then cooled to ambient temperature.

5 The mixture was poured into saturated sodium bicarbonate (80 mL) and extracted with ether (3 x 60 mL). The organic extracts were dried (magnesium sulfate), concentrated down *in vacuo* to ~ 30 mL and added to a stirred suspension of *tert*-butyl N-(1*S*,2*R*)-3-amino-1-benzyl-2-

10 hydroxypropylcarbamate (3.9 g, 14.1 mmol) in dichloroethane (30 mL) and dimethylformamide (20 mL). Sodium triacetoxyborohydride (4.5 g, 21.2 mmol) was added followed by glacial acetic acid (0.8 mL) and the reaction was stirred at ambient temperature for 16 hours.

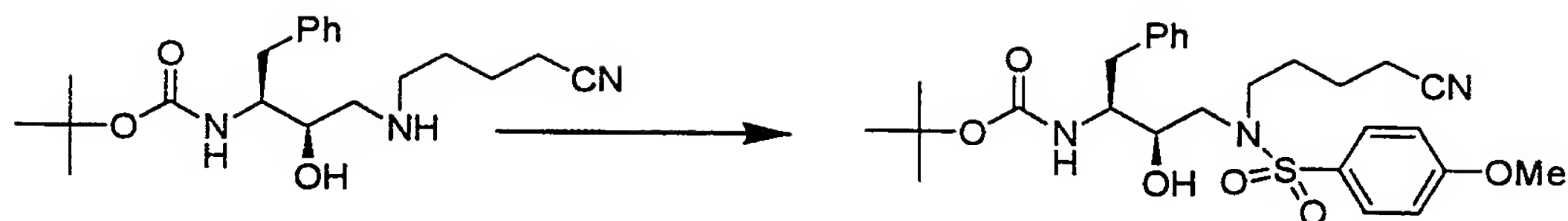
15 The mixture was filtered and the filtrate was concentrated *in vacuo* to ~25 mL, diluted with water (60 mL), adjusted to pH~12 with 1N sodium hydroxide and extracted with ethyl acetate (80 mL). The ethyl acetate was washed with water (2 x 60 mL), dried (magnesium

20 sulfate), concentrated *in vacuo* and purified by silica gel chromatography (92:8:1; chloroform: methanol: ammonium hydroxide) to afford the title compound (1.8 g, 34%) as a white solid.

¹H NMR (d₆-DMSO): δ 1.32 (9H, s), 1.48-1.72 (4H, m), 2.02 (1H, br s), 2.44-2.70 (7H, m), 3.04 (1H, dd), 3.42-3.64 (2H, m), 4.82 (1H, br s), 6.72 (1H, d), 7.15-7.35 (5H, m); MS: 362 (MH⁺).

Step 3:

-98-



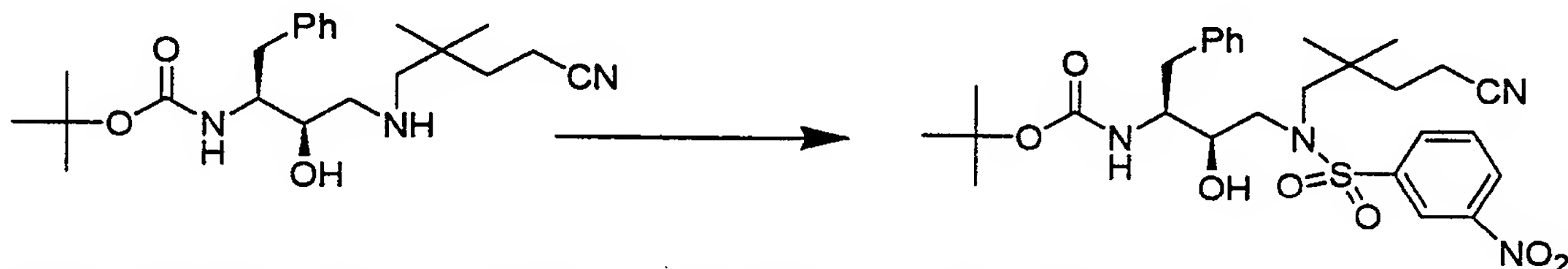
***tert*-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-(4-cyanobutyl)-2-hydroxypropyl)-4-methoxybenzenesulfonamide**

The product from Step 2 was subjected to the general procedure Step 3, Example (Compound 1) to afford the title compound as a white foam;

¹H NMR(DMSO-*d*₆): δ 1.22 (9H, s), 1.4-1.6 (4H, m), 2.4-2.5 (3H, m), 2.75 (1H, dd), 2.95 (2H, br d), 3.2-3.3 (2H, m), 3.4-3.6 (2H, m), 3.8 (3H, s), 5.07 (1H, br s), 6.7 (1H, d), 7.09 (2H, d), 7.10-7.14 (5H, m), 7.7 (1H, m); MS: 532.2 (MH⁺)

Example (Compound 5)

Step 1:



***tert*-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)-2-hydroxypropyl)-3-nitrobenzenesulfonamide**

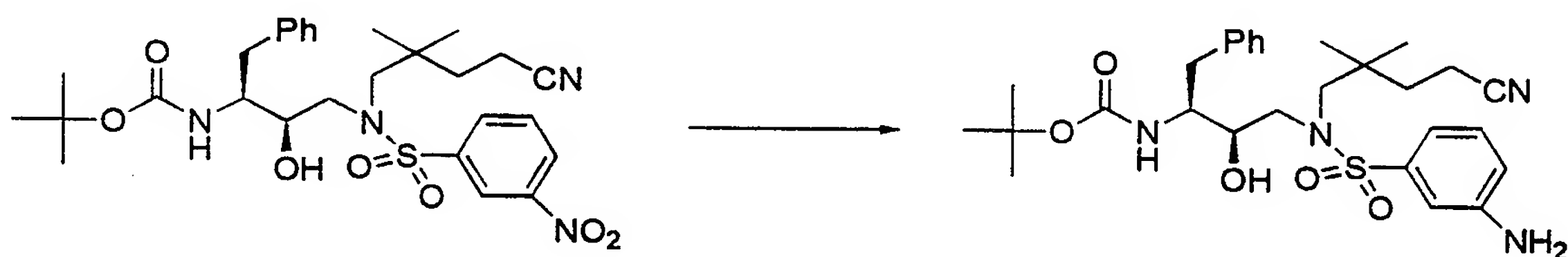
To a solution of the product from step 2 (Example (Compound 1); 0.65 g, 1.6 mmol) in dichloromethane (5 mL) was added 3-nitrobenzenesulfonyl chloride (0.4 g, 1.8 mmol) and 1M sodium hydroxide/water (3 mL, 3 mmol) and the mixture was stirred at ambient temperature for 7 hours. Excess 3-nitrobenzenesulfonyl chloride (0.2 g, 0.9 mmol) and 1M sodium hydroxide/water (1.5 mL, 1.5 mmol) were added and stirring was continued

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for 18 hours. The organic phase was dried (sodium sulfate) and evaporated. The residue was chromatographed (silica gel, dichloromethane/ methanol, 97:3) to provide the title compound (0.52 g, 55%) as a white foam;

5 ^1H NMR ($\text{DMSO}-d_6$): δ 0.9 (3H, s), 0.95 (3H, s), 1.2 (9H, s), 1.65 (2H, t), 2.4 (1H, dd), 2.50-2.55 (1H, m), 2.8 (1H, d), 3.0 (1H, d), 3.2 (1H, dd), 3.3-3.5 (5H, m), 4.95 (1H, d), 6.6 (1H, d), 7.10-7.25 (5H, m), 7.85 (1H, t), 8.15 (1H, d), 8.45 (1H, d), 8.5 (1H, s).

10 Step 2:

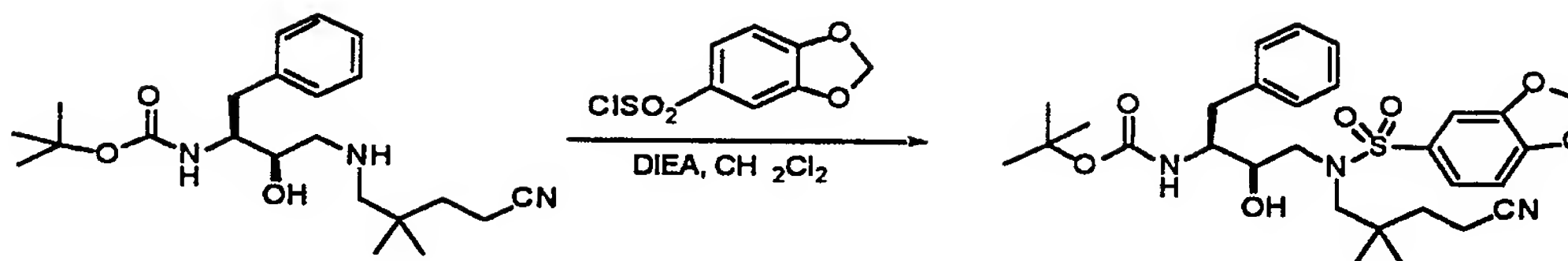


tert-ButylN-((1S,2R)-3-[(3-aminophenyl)sulfonyl](4-cyano-2,2-dimethylbutyl)amino-1-benzyl-2-hydroxypropyl)carbamate

15 A mixture of the product from step 1 (0.19 g), 10% Pd/C (40 mg), and ethylacetate (4 mL) was hydrogenated at atmospheric pressure for 18 hours. Solvent was evaporated and the residue was chromatographed (silica gel, dichloromethane/methanol, 20 97:3) to give the title compound (0.16 g, 85%) as a foam; ^1H NMR (CDCl_3): δ 0.99 (6H, s), 1.25 (9H, s), 1.75 (2H, t), 2.4 (2H, t), 2.8 (1H, dd), 2.95-3.20 (5H, m), 3.6-3.8 (2H, br m), 3.9-4.1 (3H, br m), 4.58 (1H, br d), 6.85 (1H, d), 7.05 (1H, br s), 7.09 (1H, d), 7.18-7.31 (6H, 25 m); MS: 545.2 (MH^+).

Example (Compound 6)

-100-

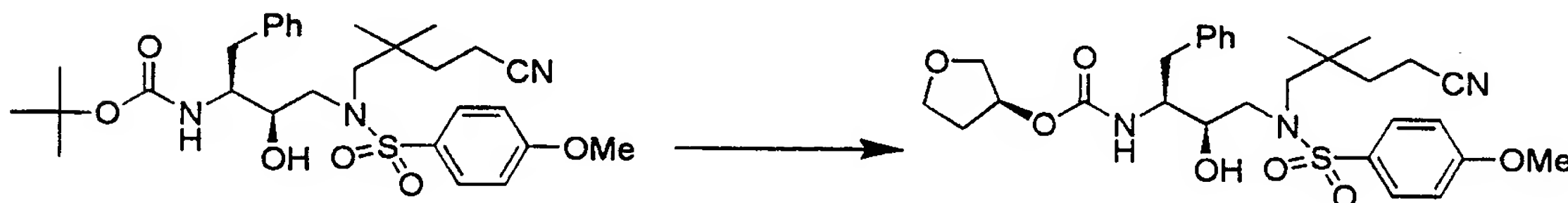


{ (1S,2R)-3-[(Benzo[1,3]dioxole-5-sulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid *tert*-butyl ester.

[(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-carbamic acid *tert*-butyl ester (0.099 g, 0.25 mmol) was dissolved in CH₂Cl₂ and treated with diisopropylethylamine (0.066 ml, 0.38 mmol) and Benzo[1,3]dioxole-5-sulfonyl chloride (0.067 g, 0.31 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (1% MeOH in CH₂Cl₂) gave 0.083 g of a white solid. MS (ES): 574 (M+1).

20

Example (Compound 7)

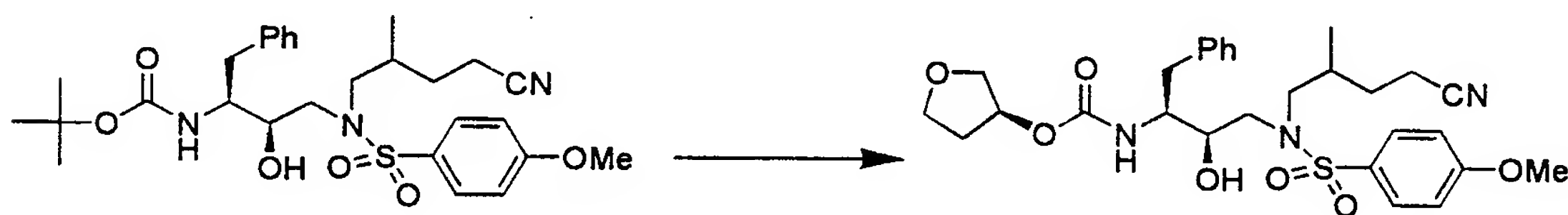


(3S)-Tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

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To a solution of the product of Step 3, Example (Compound 1) (0.22 g, 0.4 mmol) in dichloromethane (2.5 mL) at 0 °C was added trifluoroacetic acid (2.5 mL) and the mixture was stirred at ambient temperature for 1 hour. Solvent was evaporated and coevaporated with dichloromethane. The residue was dissolved in dichloromethane and washed with 0.1 M sodium hydroxide/water, water, brine, dried (sodium sulfate), concentrated and dried in vacuo. The resulting pale yellow foam was dissolved in acetonitrile (4 mL) and *N*-ethyldiisopropylamine (0.08 mL, 0.059 g, 0.46 mmol) and *N*-succinimidyl (3*S*)-tetrahydro-3-furanyl carbonate (0.098 g, 0.43 mmol) were added. The mixture was stirred at ambient temperature for 1 hour and the solvent was evaporated. The residue was chromatographed (silica gel, dichloromethane/methanol, 98:2) to give the title compound as a white foam (0.18 g, 82%);

¹H NMR (DMSO-*d*₆): δ 0.88 (3H, s), 0.92 (3H, s), 1.65 (2H, t), 1.75 (1H, dt), 2.05 (1H, sextuplet), 2.45-2.50 (2H, m), 2.70 (1H, s), 2.85 (1H, dd), 2.93 (1H, dd), 3.2-3.4 (5H, m), 3.57 (1H, dd), 3.63 (1H, td), 3.69 (1H, dd), 3.72-3.80 (1H, s), 3.84 (3H, s), 4.90 (1H, br s), 5.18 (1H, d), 7.08 (2H, d), 7.10-7.25 (6H, m), 7.75 (2H, d); MS: 574.2 (MH⁺).



Example (Compound 8)

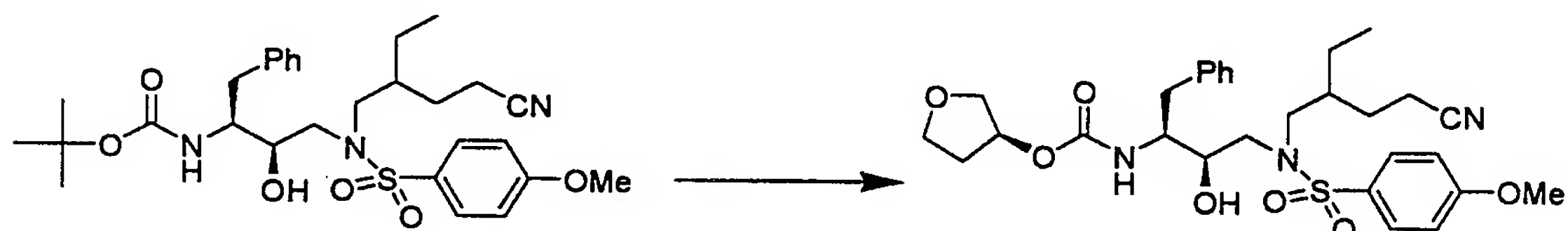
-102-

(3S)-Tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-[(2RS)-4-cyano-2-methyl][(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

The product from step 3, Example (Compound 2) was subjected to general procedure in Example (Compound 7) to give the title compound as a white glass (~1:1 mixture of diastereomers by ^1H NMR spectroscopy); ^1H NMR (DMSO- d_6): δ 0.80 (3H, d), 1.2-1.3 (1H, m), 1.75 (1H, dd), 1.8-1.9 (2H, m), 2.0-2.1 (1H, m), 2.40 (1H, dd), 2.45-2.50 (2H, m), 2.6-2.7 (2H, m), 2.98 (1H, d), 3.10 (1H, dd), 3.3-3.4 (2H, m), 3.5-3.7 (5H, m), 3.82 (3H, s), 4.90 (1H, br s), 5.10 (1H, d), 7.08 (2H, d), 7.10-7.25 (6H, m), 7.72 (2H, d); MS: 560.2 (MH^+).

15

Example (Compound 9)



(3S)-Tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-[(2RS)-4-cyano-2-ethyl][(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

20

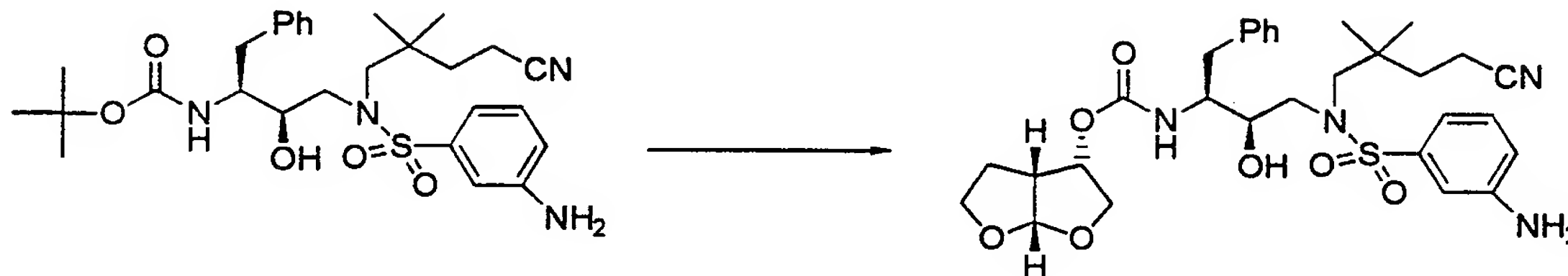
The product from Step 3, Example (Compound 3) was subjected to general procedure in Example (Compound 7) to provide the title compound as a white glass (~1:1 mixture of diastereomers by ^1H NMR spectroscopy);

^1H NMR (DMSO- d_6): δ 0.80 (3H, t), 1.2-1.4 (2H, m), 1.42-1.60 (2H, m), 1.70-1.80 (2H, m), 1.95-2.05 (1H, m), 2.4-2.5 (2H, m), 2.90-2.95 (3H, m), 2.98 (1H, dd), 3.10 (1H, dd), 3.2-3.3 (1H, m), 3.4-3.90 (6H, m), 3.94 (3H, s),

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4.90 (1H, br s), 5.05 (1H, d), 7.05-7.30 (8H, m), 7.75 (2H, d); MS: 574.2 (MH⁺).

Example (Compound 10)



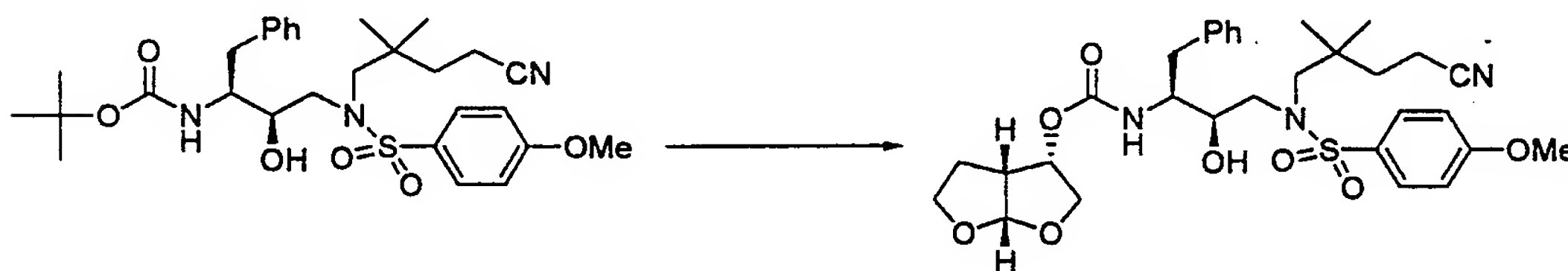
(3*S*, 3*aR*, 6*aS*) -Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-3-
5 [(3-aminophenyl)sulfonyl](4-cyano-2,2-dimethylbutyl)-
amino-1-benzyl-2-hydroxypropyl) carbamate

To a solution of the product from Step 2,
Example (Compound 5) (0.13 g) in dichloromethane (1.5 mL)
at 0°C was added trifluoroacetic acid (1 mL) and the
10 mixture was stirred at ambient temperature for 1 hour.
Solvent was evaporated and coevaporated with
dichloromethane. The residue was dissolved in
dichloromethane and washed with 0.1 M sodium
hydroxide/water, dried (sodium sulfate), concentrated and
15 dried in vacuo. The resulting pale yellow foam was
dissolved in acetonitrile (3 mL) and *N*-
ethyldiisopropylamine (0.09 mL, 0.067 g, .52 mmol) and
[(3*S*, 3*aR*, 6*aS*)-hexahydrofuro[2,3-*b*]furan-3-yl] [4-
nitrophenyl] carbonate (0.08 g, 0.31 mmol) were added.
20 The mixture was stirred at ambient temperature under
nitrogen atmosphere for 18 hours. Solvent was evaporated
and the residue was dissolved in dichloromethane, washed
with saturated sodium bicarbonate/water, dried (sodium
sulfate) and chromatographed (silica gel,
25 dichloromethane/methanol, 97:3) to give the title
compound (0.09 g, 65%) as a foam;

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^1H NMR ($\text{DMSO}-d_6$): δ 0.9 (3H, s), 0.97 (3H, s), 1.6-1.7 (3H, m), 1.8 (1H, dd), 2.2-2.3 (3H, m), 2.7 (1H, d), 2.80-2.95 (3H, m), 3.3-3.4 (3H, m), 3.45 (1H br q), 3.6-3.8 (4H, m), 4.8 (1H, q), 5.15 (1H, d), 5.58 (1H, d), 5.6 (2H, br s), 6.8 (1H, d), 6.85 (1H, d), 6.98 (1H, br s), 7.10-7.25 (7H, m); MS: 601.2 (MH^+).

Example (Compound 11)

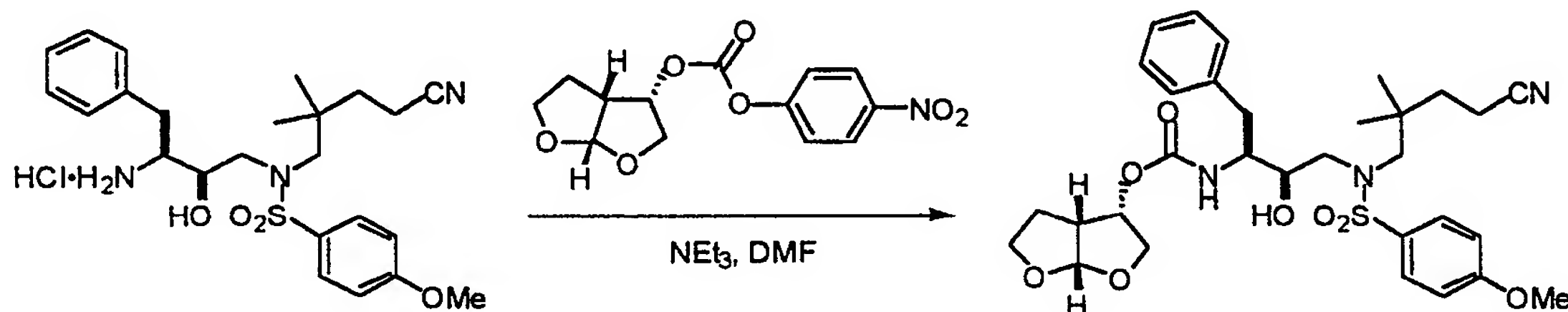


(3*S*,3*aR*,6*aS*)-Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl) [(4-methoxyphenyl) sulfonyl]amino-2-hydroxypropyl) carbamate

The product from Step 3 of Example (Compound 1) was subjected to the procedure used in Example (Compound 10) to provide the title compound as a white foam;

^1H NMR ($\text{DMSO}-d_6$): δ 0.89 (3H, s), 0.92 (3H, s), 1.6-1.7 (3H, m), 1.5 (1H, dd), 2.4-2.5 (3H, m), 2.7 (1H, d), 2.85-2.95 (3H, m), 3.25-3.45 (4H, m), 3.6-3.8 (4H, m), 3.82 (3H, s), 4.79 (1H, q), 5.15 (1H, d), 5.53 (1H, d), 7.08 (2H, d), 7.12-7.28 (6H, m), 7.75 (2H, d); MS: 616.5 (MH^+).

Alternative Synthesis



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(3*S*, 3*aR*, 6*aS*) -Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate
N-(2*R*, 3*S*)-(3-Amino-2-hydroxy-4-phenyl-butyl)-*N*-(4-cyano-2,2-dimethyl-butyl)-4-methoxy-benzenesulfonamide
 5 hydrochloride (0.086 mmol) was dissolved in 1 mL of DMF and treated with triethylamine (60 μ L, 0.43 mmol) and (3*S*, 3*aR*, 6*aS*) hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl carbonate (0.028 g, 0.095 mmol) at ambient
 10 temperature under argon with stirring. After 3 days the reaction mixture was dissolved in EtOAc, washed with water and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (EtOAc, 50% in Hexane) gave 0.028 g
 15 of a white solid. MS (ES): 616.3 (*M*+1), HPLC showed the material to be 98% pure; ret. time = 11.80 min.

Example (Compound 12)

Step 1:



tert-Butyl *N*-[(1*S*, 2*R*)-1-benzyl-3-((4-cyano-2,2-dimethylbutyl)[3-(methylamino)phenyl]sulfonylamino)-2-hydroxypropyl]carbamate
 20

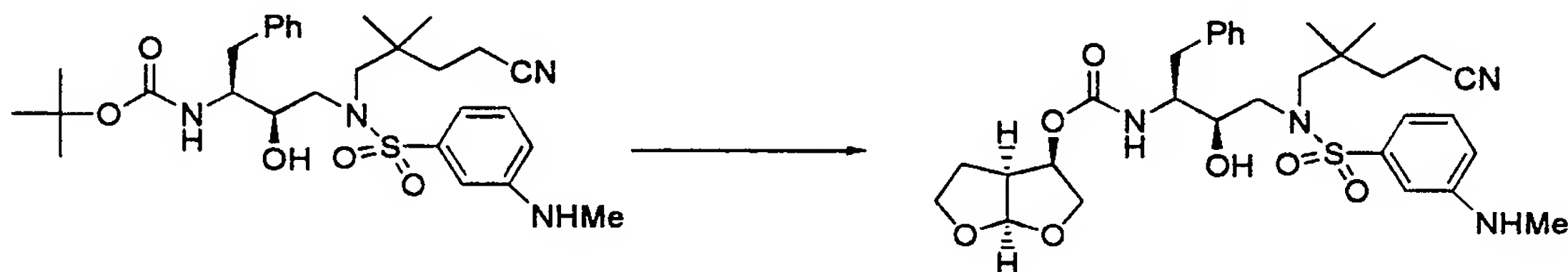
A mixture of the product from Step 1 (Example (Compound 5), 0.21 g), 10% Pd/C (0.2 g), methanol (7 mL), and acetic acid (0.3 mL) was hydrogenated at atmospheric

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pressure for 48 hours. The mixture was filtered through a bed of celite and the solvent was evaporated. The residue was dissolved in dichloromethane and the solution was washed with saturated sodium bicarbonate/water, dried (sodium sulfate), evaporated, and the residue was chromatographed (silica gel, dichloromethane/methanol, 98:2) to afford the title compound (25 mg, 13%) as a glass;

¹H NMR (CDCl₃): δ 0.95 (6H, s), 1.29 (9H, s), 1.72 (2H, t), 2.37 (2H, t), 2.75-2.85 (1H, m), 2.87 (3H, s), 2.90-3.20 (5H, m), 3.75 (1H, brs), 3.95 (1H, brs), 4.02 (1H, brs), 4.18 (1H, brs), 4.55 (1H, d), 6.76 (1H, dd), 6.95 (1H, t), 7.02 (1H, d), 7.18-7.23 (3H, m), 7.25-7.32 (3H, m).

Step 2:



(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-benzyl-3-((4-cyano-2,2-dimethylbutyl)[3-(methylamino)-phenyl]sulfonylamino)-2-hydroxypropyl]carbamate

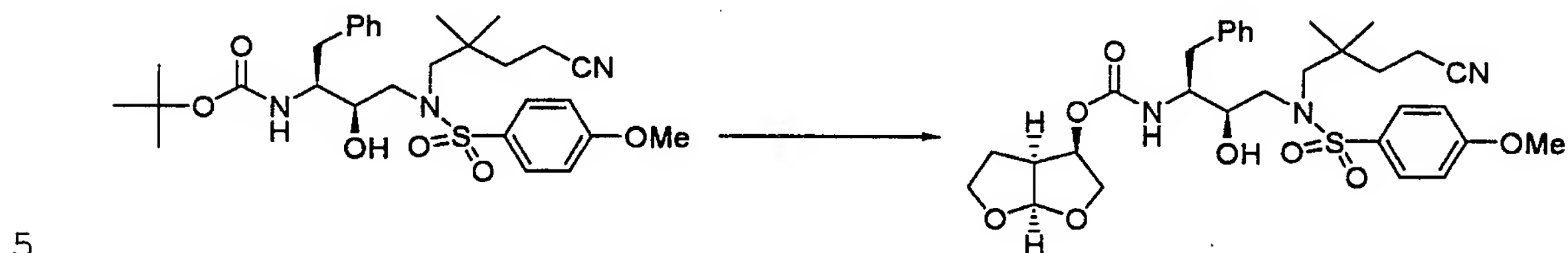
The product from Step 1 was subjected to the procedure used in Example (Compound 10) to afford the title compound as a white foam;

¹H NMR (DMSO-d₆): δ 0.97 (3H, s), 1.02 (3H, s), 1.20-1.26 (1H, m), 1.30-1.35 (1H, m), 1.78 (2H, t), 2.50-2.62 (3H, m), 2.78 (3H, d), 2.80-2.97 (3H, m), 3.05 (1H, d), 3.38-3.42 (2H, m), 3.50-3.60 (1H, m), 3.65 (2H, dd), 3.78 (1H,

-107-

td), 3.90 (2H, dd), 4.85 (1H, q), 5.26 (1H, d), 5.58 (1H, d), 6.24 (1H, brq), 6.85 (1H, d), 6.94-7.0 (2H, m), 7.2-7.4 (7H, m); MS: 615.5 (MH⁺).

Example (Compound 13)

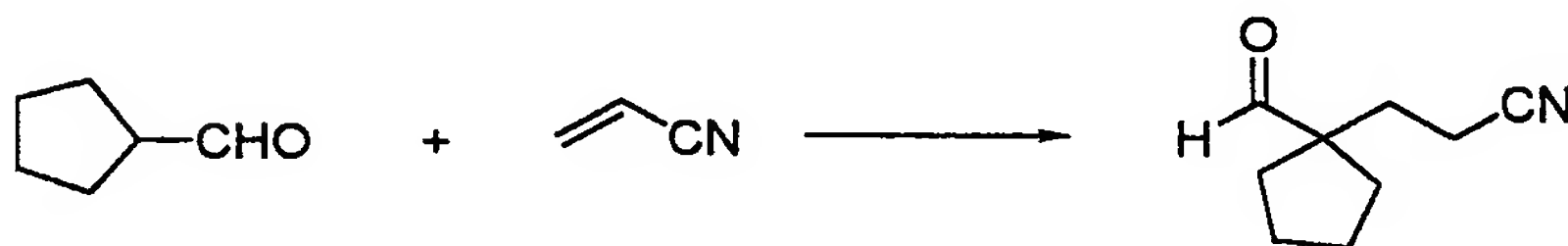


(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl) [(4-methoxyphenyl) sulfonyl]amino-2-hydroxypropyl) carbamate

The product from Step 3 (Example (Compound 1)) was subjected to the procedure used in Example (Compound 11), except the enantiomeric activated ester was employed, to afford the title compound as a white foam; ¹H NMR (DMSO-*d*₆): δ 0.92 (3H, s), 1.0 (3H, s), 1.25 (1H, d), 1.38-1.48 (1H, m), 1.79 (2H, t), 2.43-2.60 (3H, m), 2.75 (1H, d), 2.8 (1H, dd), 2.9 (1H, dd), 3.05 (1H, dd), 3.3-3.5 (4H, m), 3.6-3.8 (4H, m), 3.92 (3H, s), 4.86 (1H, q), 5.25 (1H, d), 5.6 (1H, d), 7.18 (2H, d), 7.22-7.40 (6H, m), 7.8 (2H, d); MS: 616.4 (MH⁺).

Example (Compound 14)

20 Step 1:



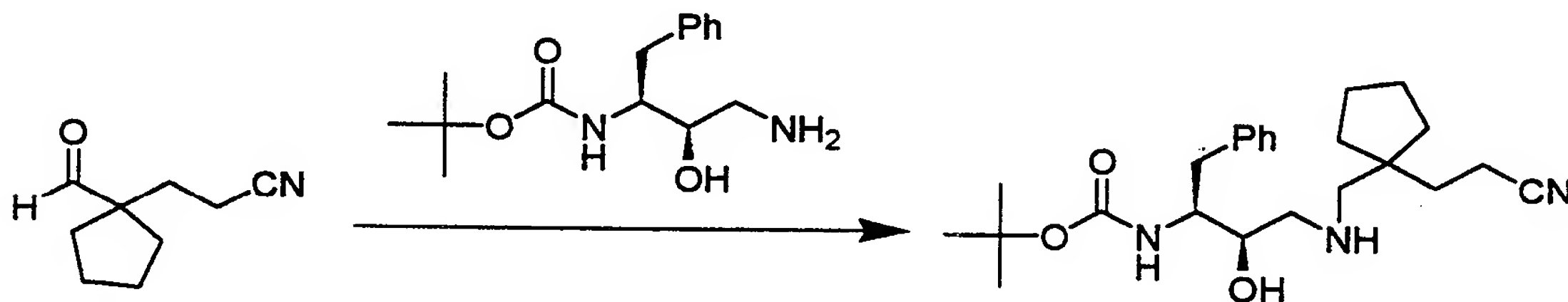
3-(1-Formylcyclopentyl)propanenitrile

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Cyclopentanecarboxaldehyde was subjected to Step 1 of Example (Compound 1) to provide the title compound.

¹H NMR (CDCl₃): δ 1.44-1.58 (2H, m), 1.63-1.80 (4H, m),
5 1.89-2.01 (4H, m), 2.29 (2H, t), 9.40 (1H, s).

Step 2:

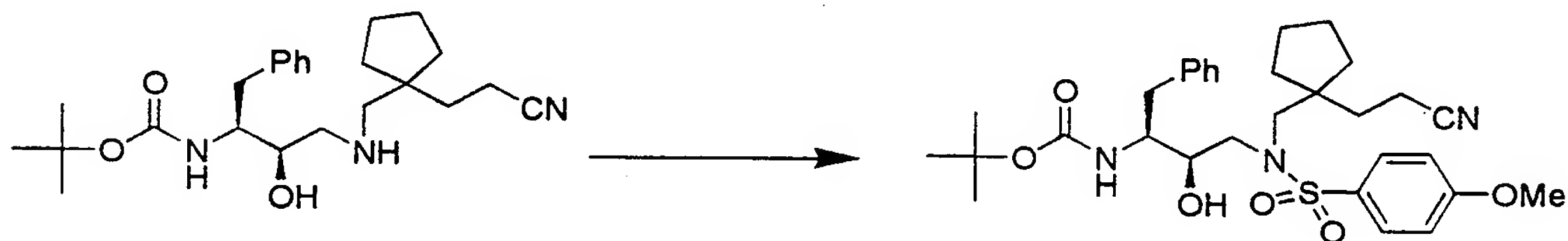


tert-Butyl N-[(1*S*,2*R*)-1-benzyl-3-([1-(2-cyanoethyl)
cyclopentyl]methylamino)-2-hydroxypropyl]carbamate

10 The product from Step 1 was subjected to procedure 2 (Example (Compound 1)) to give the title compound as a thick yellow oil.

¹H NMR (DMSO-*d*₆): δ 1.24 (9H, s), 1.28-1.60 (8H, m), 1.64
(2H, t), 2.29 (2H, s), 2.40 (2H, t), 2.46-2.62 (3H, m),
15 2.96 (1H, dd), 3.36-3.58 (2H, m), 4.75 (1H, br s), 6.70
(1H, d), 7.08-7.25 (5H, m); MS 416 (MH⁺).

Step 3:



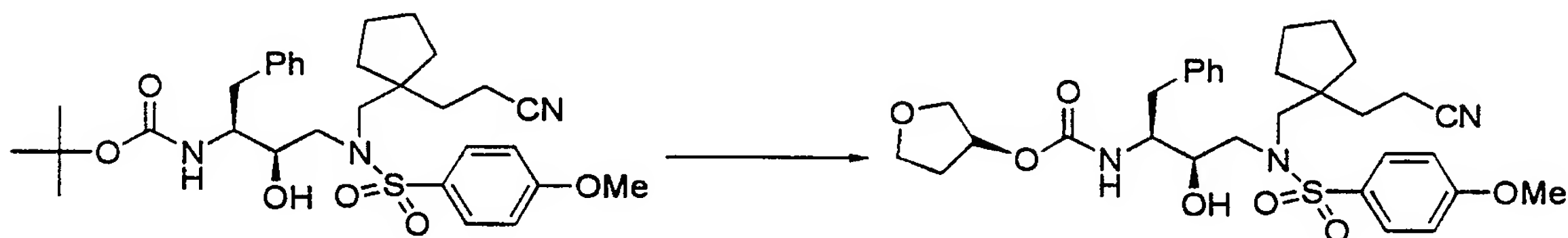
tert-Butyl N-((1*S*,2*R*)-1-benzyl-3-[1-(2-cyanoethyl)
cyclopentyl]methyl[(4-methoxyphenyl)sulfonyl]amino-2-
20 hydroxypropyl]carbamate

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The product from Step 2 was subjected to Step 3 (Example (Compound 1)) to afford the title compound as a white solid.

¹H NMR (DMSO-*d*₆): δ 1.22 (9H, s), 1.38-1.60 (8H, m), 1.66-1.76 (1H, m), 1.82-1.92 (1H, m), 2.44-2.56 (2H, m), 2.68-2.96 (3H, m), 3.24-3.44 (4H, m), 3.68-3.78 (1H, m), 3.82 (3H, s), 5.08 (1H, d), 6.64 (1H, d), 7.04-7.24 (7H, m), 7.74 (2H, d); MS: 586 (MH⁺).

Step 4:



(3*S*)-Tetrahydro-3-furanyl-*N*-((1*S*,2*R*)-1-benzyl-3-[1-(2-cyanoethyl)cyclopentyl]methyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

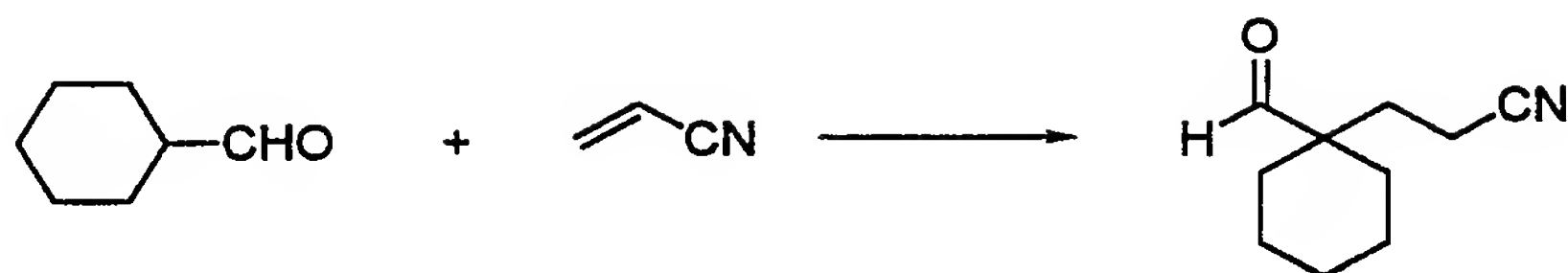
The product from Step 3 was subjected to the procedure used in Example (Compound 7) to give the title compound as a white foam.

¹H NMR (DMSO-*d*₆): δ 1.4-1.7 (8H, m), 1.75-2.00 (3H, m), 2.1 (1H, sextuplet), 2.5-2.6 (2H, m), 2.8 (1H, d), 2.9 (1H, dd), 3.0 (1H, d), 3.3-3.5 (5H, m), 3.65 (1H, dd), 3.7 (1H, dd), 3.8 (1H, dd), 3.82-3.87 (1H, m), 3.92 (3H, s), 5.0 1H, brs), 5.21 (1H, d), 7.18 (2H, d), 7.2-7.4 (6H, m), 7.82 (2H, d); MS: 600.4 (MH⁺).

Example (Compound 15)

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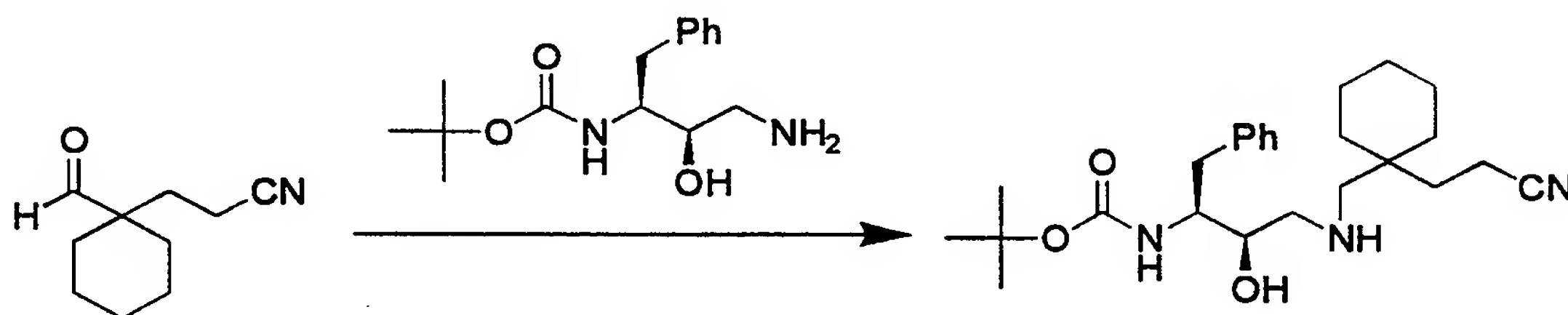
Step 1:

**3-(1-Formylcyclohexyl)propanenitrile**

Cyclohexylcarboxaldehyde was subjected to Step 5 in Step 1 of Example (Compound 1) to give the title compound.

¹H NMR (CDCl₃): δ 1.22-1.43 (5H, m), 1.47-1.62 (3H, m), 1.81-1.91 (4H, m), 2.21 (2H, t), 9.40 (1H, s)

Step 2:



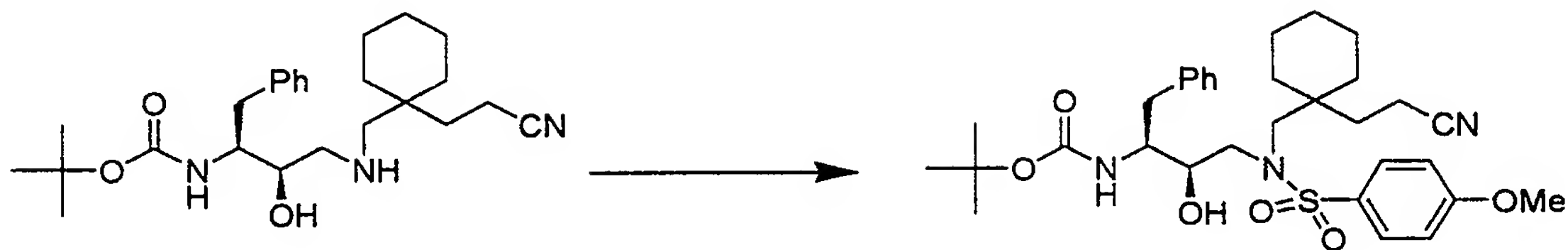
10 tert-Butyl N-[(1S,2R)-1-benzyl-3-([1-(2-cyanoethyl)cyclohexyl]methylamino)-2-hydroxypropyl]carbamate

The product from step 1 was subjected to Step 2 (Example (Compound 1)) to afford the title compound as a thick yellow oil.

15 ¹H NMR (DMSO-d₆): δ 1.24 (9H, s), 1.18-1.42 (10H, m), 1.60 (2H, t), 2.30 (2H, s), 2.39 (2H, t), 2.46-2.62 (3H, m), 2.96 (1H, dd), 3.36-3.58 (2H, m), 4.74 (1H, br s), 6.71 (1H, d), 7.08-7.26 (5H, m); MS: 430 (MH⁺).

Step 3:

-111-

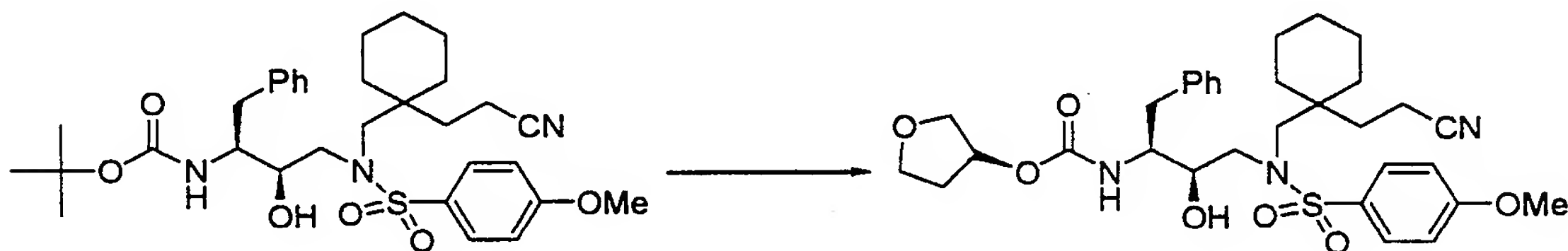


tert-Butyl N-((1S,2R)-1-benzyl-3-[1-(2-cyanoethyl)cyclohexyl]methyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

5 The product from Step 2 was subjected to Step 3 (Example (Compound 1)) to provide the title compound as a white solid.

¹H NMR (DMSO-d₆): δ 1.20 (9H, s), 1.18-1.52 (10H, m), 1.75-1.90 (2H, m), 2.40-2.50 (2H, m), 2.64-2.94 (3H, m),
 10 3.22-3.38 (4H, m), 3.70-3.78 (1H, m), 3.82 (3H, s), 5.11 (1H, d), 6.64 (1H, d), 7.04-7.24 (7H, m), 7.76 (2H, d);
 MS: 600 (MH⁺)

Step 4:



15 **(3S)-Tetrahydro-3-furanyl-N-((1S,2R)-1-benzyl-3-[1-(2-cyanoethyl)cyclohexyl]methyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate**

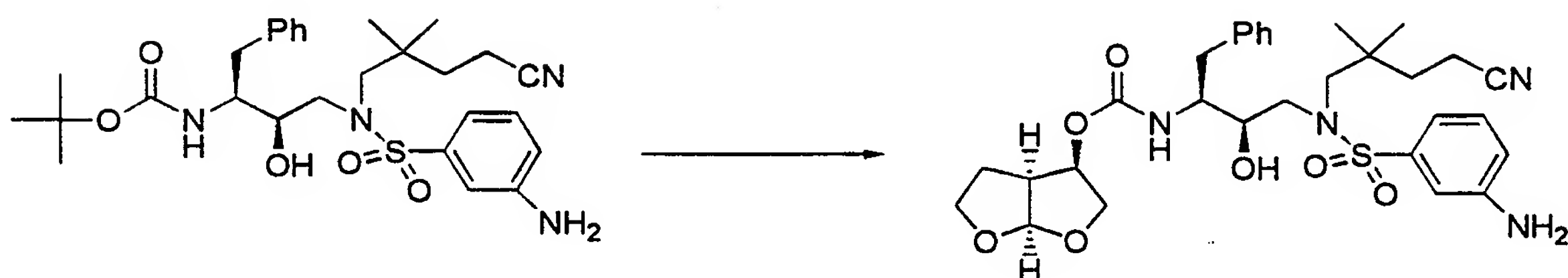
The product from Step 3 was subjected to conditions used in Example (Compound 7) to give the title
 20 compound as a white foam.

¹H NMR (DMSO-d₆): δ 1.2-1.6 (10H, m), 1.7-1.9 (3H, m), 2.05 (1H, sextuplet), 2.4-2.5 (2H, m), 2.65 (1H, d), 2.8 (1H, dd), 2.92 (1H, d), 3.2-3.4 (5H, m), 3.55 (1H, dd), 3.65 (1H, dd), 3.72 (1H, dd), 3.80 (1H, brs), 3.85 (3H,

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s), 4.9 (1H, brs), 5.10 (1H, d), 7.08 (2H, d), 7.1-7.3 (6H, m), 7.8 (2H, d); MS: 614.4 (MH⁺).

Example (Compound 16)



(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-
5 [(3-aminophenyl)sulfonyl](4-cyano-2,2-dimethylbutyl)
amino-1-benzyl-2-hydroxypropyl) carbamate

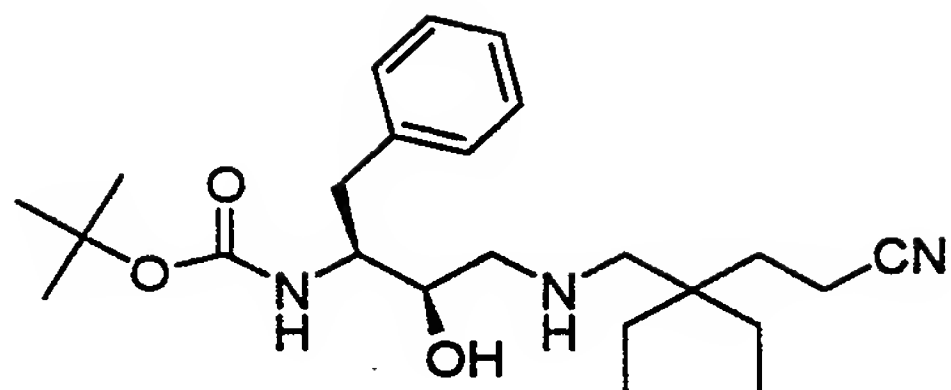
The product from Example (Compound 5) was
subjected to general procedure in Example (Compound 11)
using [(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl][4-
10 nitrophenyl] carbonate to give the title compound as a
white foam.

¹H NMR (DMSO-d₆): δ 0.95 (3H, s), 0.98 (3H, s), 1.15 (1H,
dd), 1.3-1.4 (1H, m), 1.7 (2H, t), 2.4-2.5 (3H, m), 2.68
(1H, d), 2.72 (1H, dd), 2.8 (1H, dd), 3.25-3.35 (2H, m),
15 3.40-3.47 (1H, m), 3.57 (1H, dd), 3.7 (1H, td), 3.8 (1H,
d), 3.83 (1H, d), 4.79 (1H, dd), 5.2 (1H, d), 5.48 (1H,
d), 5.56 (2H, s), 6.78 (1H, dd), 6.86 (1H, d), 6.97 (1H,
s), 7.10-7.25 (7H, m); MS: 601.1 (MH⁺).

Example (Compound 17)

20 Step 1:

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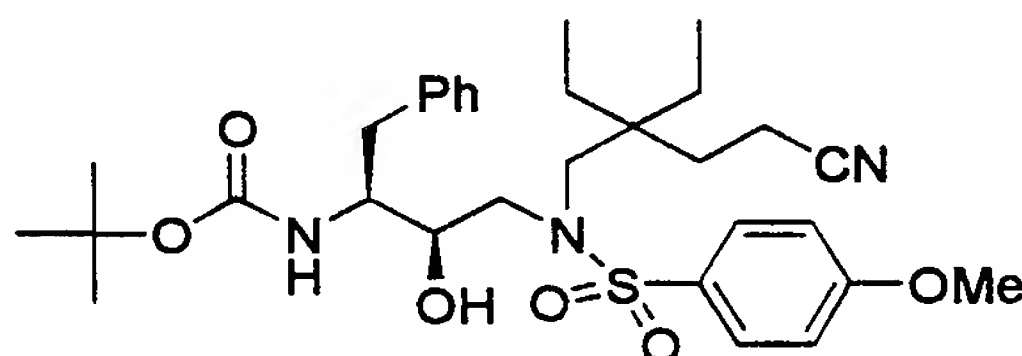


tert-Butyl N-((1S,2R)-1-benzyl-3-[(4-cyano-2,2-diethylbutyl)amino]-2-hydroxypropyl)carbamate

3,3-Diethyl-4-oxobutyl cyanide was subjected to
 5 Step 2, Example (Compound 1) to provide the title
 compound as a foam.

¹H NMR (DMSO-*d*₆): δ 0.90 (6H, t), 1.16 (4H, q), 1.24 (9H, s), 1.36 (1H, br), 1.50 (2H, t), 2.21 (2H, s), 2.36 (2H, t), 2.45-2.60 (3H, m), 2.95 (1H, dd), 3.36-3.58 (2H, m),
 10 4.74 (1H, d), 6.71 (1H, d), 7.08-7.26 (5H, m) ; MS 418 (MH⁺).

Step 2:



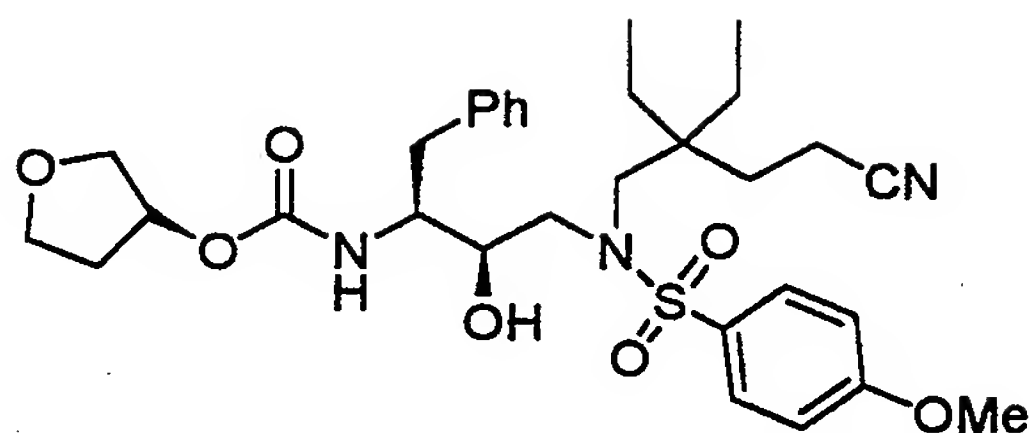
tert-Butyl N-((1S,2R)-1-benzyl-3-[(4-cyano-2,2-diethylbutyl)[(4-methoxyphenyl)sulfonyl]amino]-2-hydroxypropyl)carbamate

The product from the previous step was
 subjected to Step 3 (Example (Compound 1)) to give the
 title compound as a foam.

¹H NMR (DMSO-*d*₆): δ 0.75 (6H, dt), 1.20 (9H, s), 1.15-1.50
 20 (4H, m), 1.65-1.75 (2H, m), 2.40-2.50 (2H, m), 2.60-2.95 (3H, m), 3.22-3.42 (4H, m), 3.72-3.82 (1H, m), 3.85 (3H, s), 5.14 (1H, d), 6.62 (1H, d), 7.04-7.26 (7H, m), 7.75 (2H, d); MS 588 (MH⁺).

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Step 3:



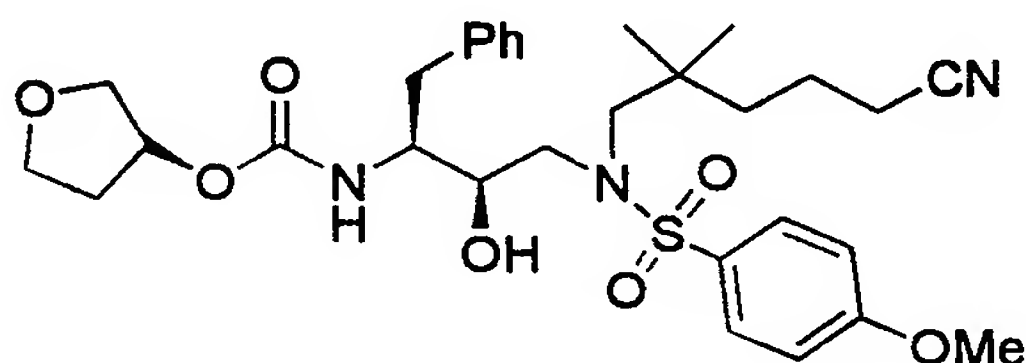
(3S)-Tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-(4-cyano-
2,2-diethylbutyl) [(4-methoxyphenyl) sulfonyl]amino-2-
5 hydroxypropyl) carbamate

The product from the previous step was subjected to the procedure used in Example (Compound 10) to give the title compound as white foam.

¹H NMR (DMSO-*d*₆): δ 0.70 (6H, s), 1.15-1.20 (3H, m), 1.35-
10 1.40 (m, 1H), 1.60-1.80 (3H, m), 2.0-2.1 (1H, m), 2.25-
2.35 (2H, m), 2.6-2.8 (2H, m), 2.90 (1H, d), 3.2-3.4 (5H, m), 3.5-3.8 (4H, m) 3.83 (3H, s), 4.90 (1H, br s), 5.2 (1H, d), 7.05 (2H, d), 7.2-7.3 (6H, m), 7.7 (2H, d); MS: 602 (MH⁺).

15

Example (Compound 18)



Step 1:

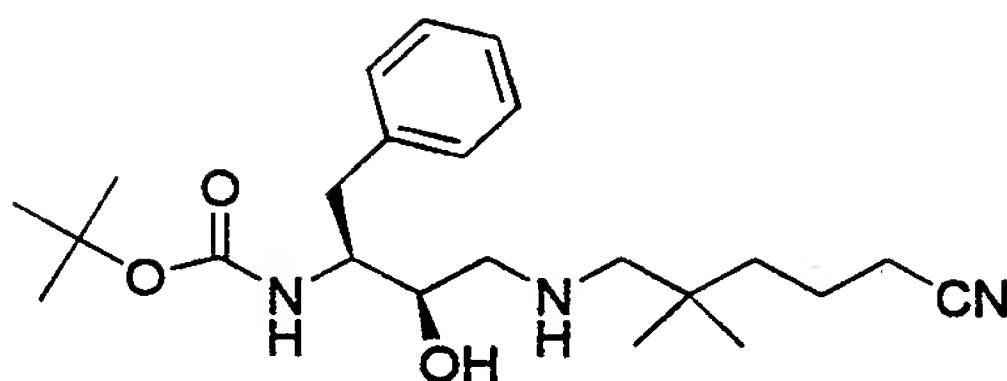
4-(1,3-Dioxolan-2-yl)-4-methylpentanenitrile

The product from Step 2 was subjected to Step 1 (Example (Compound 8) except sodium cyanide was used in
20 place of sodium azide to provide the title compound.

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^1H NMR (CDCl_3) : δ 0.92 (6H, s), 1.41-1.48 (2H, m), 1.65-1.75 (2H, m), 2.32 (2H, t), 3.82-3.97 (4H, m), 4.53 (1H, s).

Step 2:

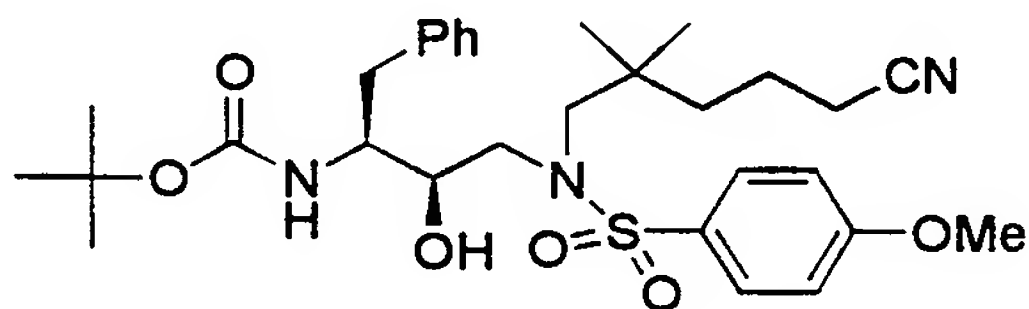


5 **tert-Butyl N-(1S,2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate**

The product from Step 1 was subjected to Step 2 (Example (Compound 8) to give the title compound as a foam.

10 ^1H NMR ($\text{DMSO}-d_6$): δ 0.82 (6H, s), 1.18-1.30 (11H, m), 1.40-1.53 (2H, m), 2.24 (2H, s), 2.38-2.62 (5H, m), 2.96 (1H, dd), 3.36-3.56 (2H, m), 4.75 (1H, br), 6.71 (1H, d), 7.08-7.26 (5H, m); MS 404 (MH+).

15 Step 3:



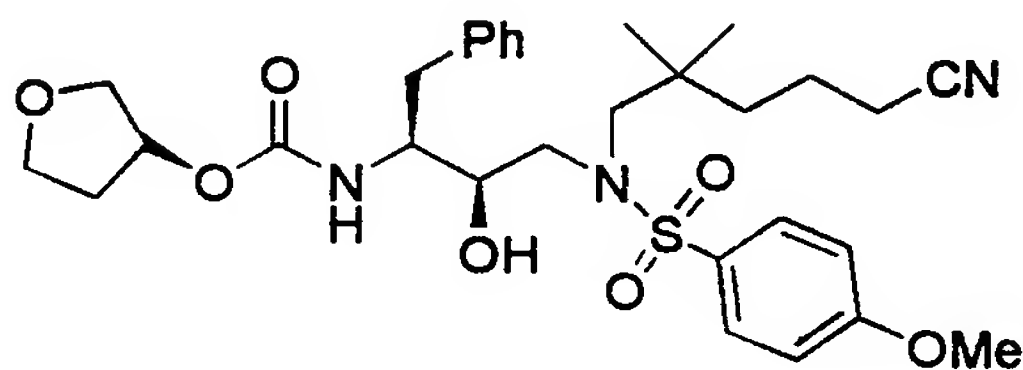
tert-Butyl N-((1S,2R)-1-benzyl-3-(5-cyano-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino)-2-hydroxypropylcarbamate

20 The product from the previous step was subjected to Step 3 (Example (Compound 1) to provide the title compound.

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¹H NMR (DMSO-*d*₆): δ 0.88 (3H, s), 0.92 (3H, s), 1.21 (9H, s), 1.26-1.37 (2H, m), 1.45-1.58 (2H, m), 2.43 (2H, t), 2.77 (1H, d), 2.87-2.97 (2H, m), 3.26-3.43 (4H, m), 3.66-3.75 (1H, m), 3.82 (3H, s), 4.98 (1H, d), 6.62 (1H, d),
 5 7.06 (2H, d), 7.08-7.24 (5H, m), 7.73 (2H, d) ; MS 574 (MH⁺) .

Step 4:



(3*S*)-Tetrahydro-3-furanyl *N*-((1*S*,2*R*)-1-benzyl-3-(5-cyano-2,2-dimethylpentyl) [(4-methoxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate
 10

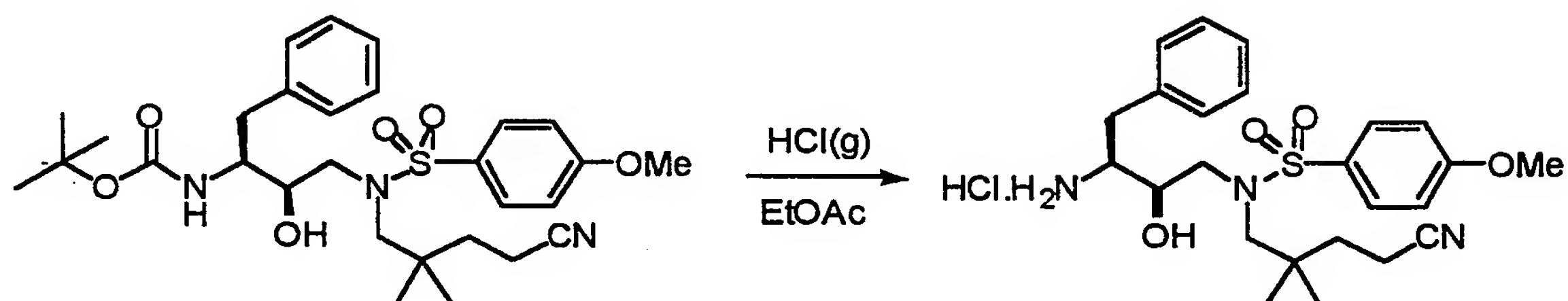
The product from the previous step was subjected to the procedure outlined in Example (Compound 7 to provide the title compound as a foam.

¹H NMR (DMSO-*d*₆): δ 0.84 (3H, s), 0.89 (3H, s), 1.3 (2H, t), 1.4-1.6 (2H, m), 1.7-1.8 (1H, m), 1.95-2.05 (1H, m), 2.4-2.5 (2H, m), 2.77 (1H, d), 2.90-2.98 (2H, br d), 3.3-3.4 (5H, m), 3.56 (1H, dd), 3.6-3.8 (3H, m), 3.82 (3H, s), 4.90 (1H, br s), 5.03 (1H, d), 7.06 (2H, d), 7.1-7.2 (6H, m), 7.7 (2H, d); MS: 588 (MH⁺) .
 15

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Example (Compound 19)

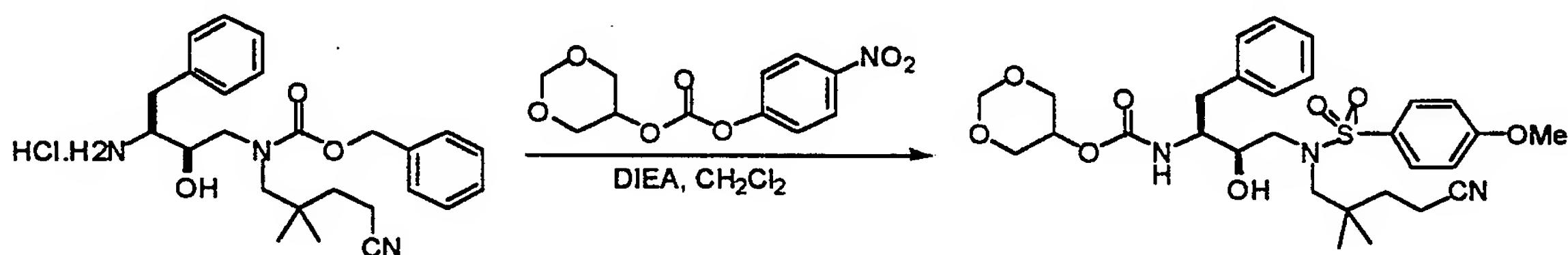
Step 1:



- 5 ***N*-(2*R*,3*S*)-(3-Amino-2-hydroxy-4-phenyl-butyl)-*N*-(4-cyano-2,2-dimethyl-butyl)-4-methoxy-benzenesulfonamide; hydrochloride.**

A solution of {(1*S*,2*R*)-1-Benzyl-3-[(4-cyano-2,2-dimethyl-butyl)-(4-methoxy-benzenesulfonyl)-amino]-2-hydroxy-propyl}-carbamic acid *tert*-butyl ester (0.600 g, 1.07 mmol) in EtOAc was cooled to -15 °C and dry HCl gas bubbled through for 30 min. The reaction was warmed to ambient temperature and solvent removed *in vacuo* to give desired product as a white solid which was used directly
10
15 in the next step.

Step 2:



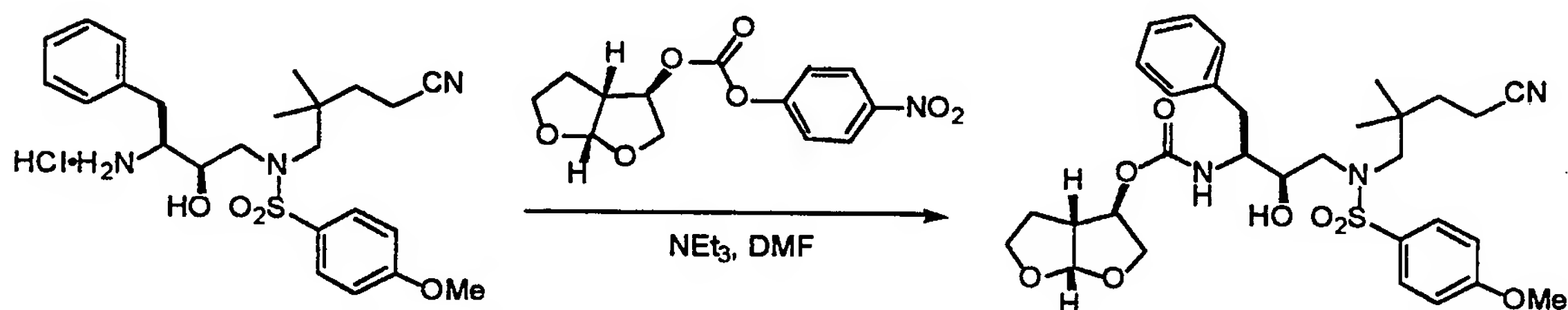
- 20 **{(1*S*,2*R*)-1-Benzyl-3-[(4-cyano-2,2-dimethyl-butyl)-(4-methoxy-benzenesulfonyl)-amino]-2-hydroxy-propyl}-carbamic acid [1,3]dioxan-5-yl ester.**

N-(2*R*,3*S*)-(3-Amino-2-hydroxy-4-phenyl-butyl)-*N*-(4-cyano-2,2-dimethyl-butyl)-4-methoxy-benzenesulfonamide
25 hydrochloride (0.111 g, 0.22 mmol) was dissolved in

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CH₂Cl₂ and treated with diisopropylethylamine (0.120 ml, 0.67 mmol) and 1,3-dioxan-5-yl 4-nitrophenyl carbonate (0.066 g, 0.25 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (1% to 3% MeOH in CH₂Cl₂) gave 0.084 g of a white solid. MS (ES): 590 (M+1).

Example (Compound 20)



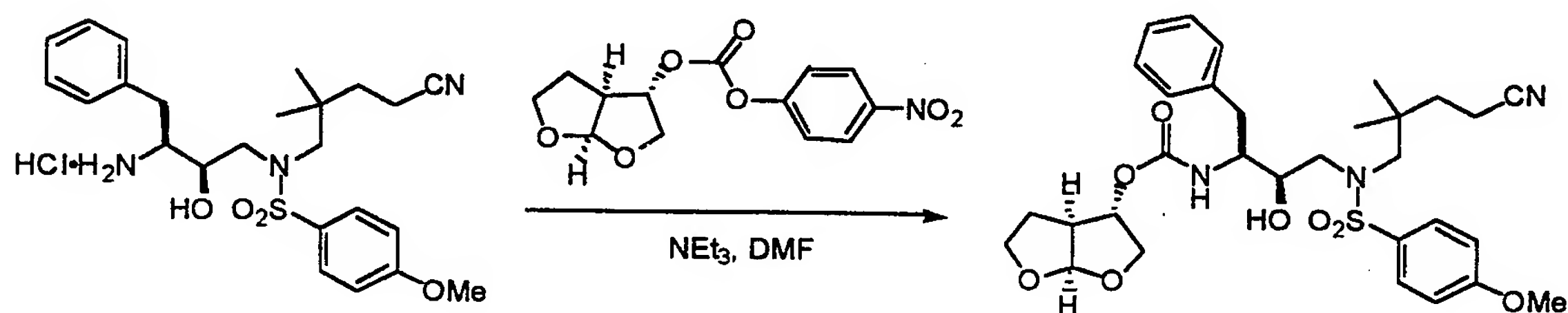
{ (1S,2R) -3- [(3-Methoxybenzenesulfonyl) - (4-cyano-2,2-dimethyl-butyl) -amino] -1-benzyl-2-hydroxy-propyl} - carbamic acid (3R,3aR,6aS)hexahydro-furo[2,3-b]furan-3-yl ester.

N- (2R,3S) - (3-Amino-2-hydroxy-4-phenyl-butyl) -N- (4-cyano-2,2-dimethyl-butyl) -4-methoxy-benzenesulfonamide hydrochloride (0.11 mmol) was dissolved in 1 mL of DMF and treated with triethylamine (75 μ L, 0.5 mmol) and (3R, 3aR, 6aS) hexahydrofuro[2,3-b]furan-2-yl 4-nitrophenyl carbonate (0.034 g, 0.12 mmol) at ambient temperature under argon with stirring. After 2 days the reaction mixture was dissolved in EtOAc, washed with water and brine. The organic phase was dried over MgSO₄, filtered

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and solvent removed *in vacuo*. Purification by column chromatography (EtOAc 50% in Hexane) gave 0.037 g of a white solid. MS (ES): 616.3 (M+1), HPLC showed the material to be 98% pure; ret. time = 11.55 min.

5

Example (Compound 21)

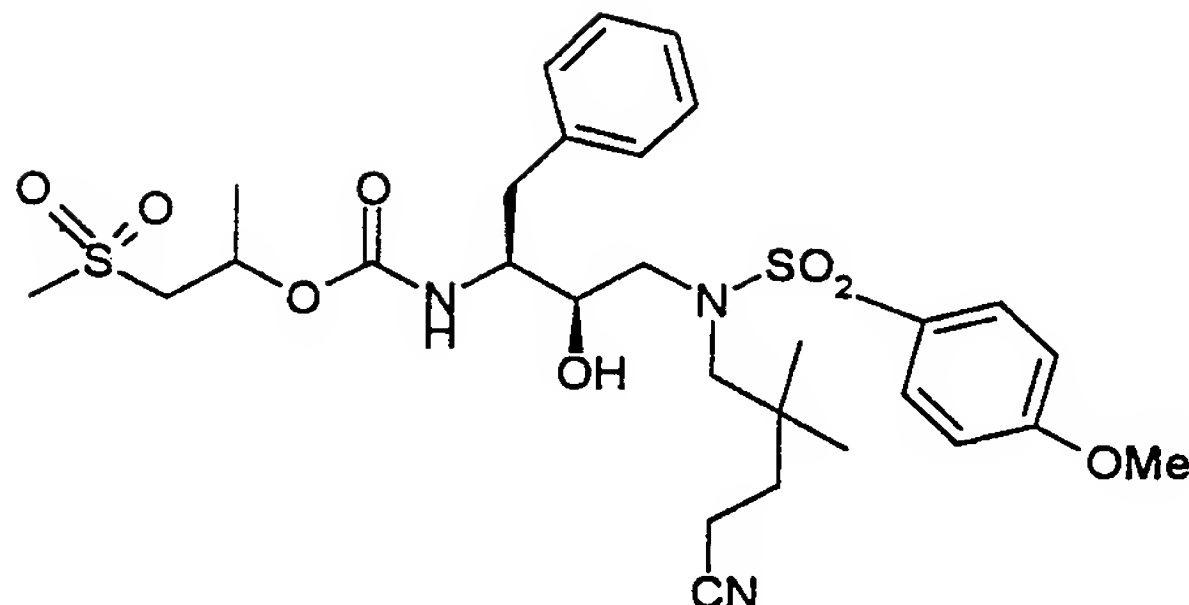
{ (1S,2R)-3-[(3-Methoxybenzenesulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (3S,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

10

N-(2R,3S)-(3-Amino-2-hydroxy-4-phenyl-butyl)-N-(4-cyano-2,2-dimethyl-butyl)-4-methoxy-benzenesulfonamide hydrochloride (0.11 mmol) was dissolved in 1 mL of DMF and treated with triethylamine (77 μ L, 0.55 mmol) and (3S, 3aS, 6aR) hexahydrofuro[2,3-b]furan-2-yl 4-nitrophenyl carbonate (0.036 g, 0.12 mmol) at ambient temperature under argon with stirring. After 2 days the reaction mixture was dissolved in EtOAc, washed with water and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (EtOAc 50% in Hexane) gave 0.044 g of a white solid. MS (ES): 616.3 (M+1), HPLC showed the material to be 98% pure; ret. time = 11.46 min.

20

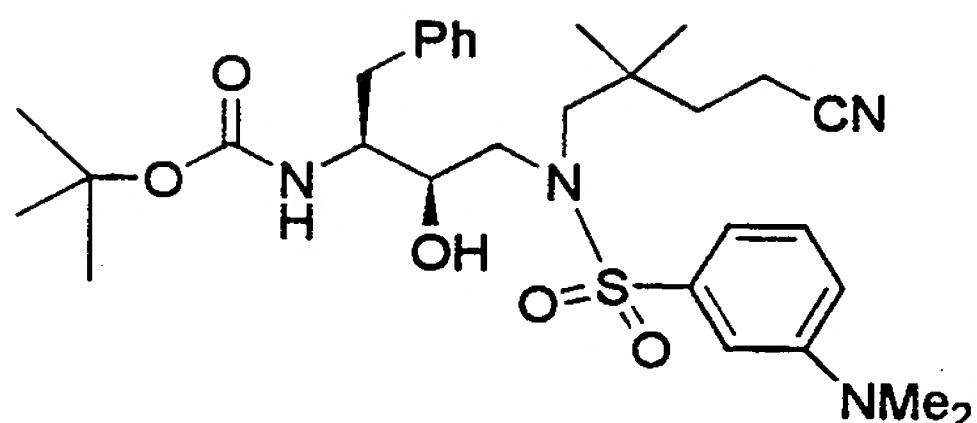
-120-

Example (Compound 22)

1N-(3-Methylsulfonylisobutyryl)-(1S,2R)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)[4-methoxyphenylsulfonyl]amino-2-hydroxypropyl amine

- 5 (1S,2R)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)
[(4-methoxy phenyl)sulfonyl]amino-2-hydroxypropylamine
(13.4, 0.027 mmol) was combined with 3-methyl
sulfonylisobutyric acid (0.007 g, 0.032 mmol) and 1-
hydroxybenzotriazole hydrate (0.004 g, 0.03 mmol) in
10 anhydrous DMF (1 ml). Triethylamine (0.020 ml) was added
followed by 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide
hydrochloride (0.007 g, 0.035 mmol). The reaction was
stirred at room temperature for 2 hours, diluted in EtOAc
and washed with sat. NaHCO₃, 0.5N KHSO₄ and brine.
- 15 Organic phase was dried with MgSO₄ and the solvent was
removed in vacuo. Purification by preparative TLC (3:1
EtOAc/Hex). Isolated 0.004 g of the product as a
colorless residue. R_f=0.40 (3:1 EtOAc/Hex LRMS (M+H)⁺
609 amu. HPLC analysis (C18); obs two peaks Tr = 10.8
20 and 11.07 min., derived from mixture of epimers at alpha
methyl proprionate linkage.

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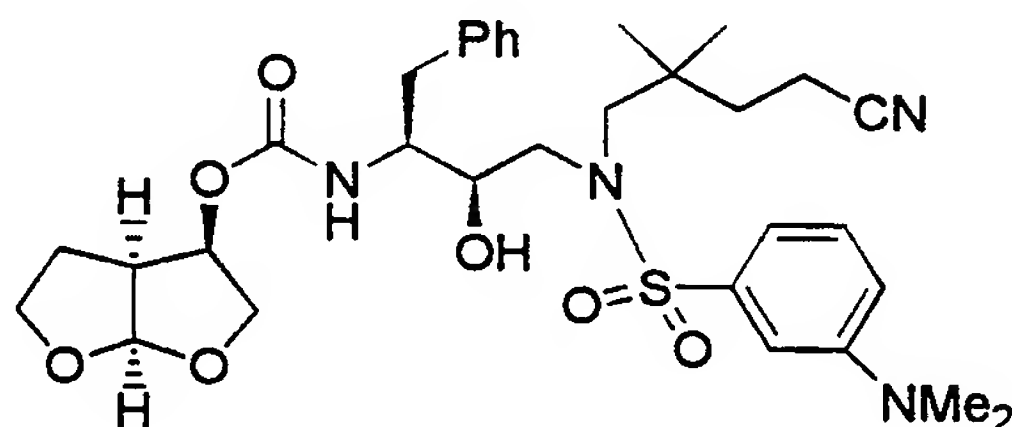
Example (Compound 23)

tert-Butyl N-((1S,2R)-3-(4-cyano-2,2-dimethylbutyl) [(3-dimethylaminophenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate

The product from Example (Compound 5) (0.10g) in ethanol (3 mL) and 37% HCOH/water (0.4 mL) was hydrogenated at atmospheric pressure with 10% Pd/C (25 mg) for 18 hours. The mixture was filtered through a bed celite, evaporated, and dissolved in dichloromethane. The solution was washed with water, dried (sodium sulfate), evaporated, and dried in vacuo to provide the title compound (90 mg) as a foam.

¹H NMR (DMSO-d₆): δ 0.89 (3H, s), 0.93 (3H, s), 1.2 (9H, s), 1.7 (2H, t), 2.4-2.5 (3H, m), 2.7 (1H, d), 2.8-2.9 (2H, m), 2.96 (6H, s), 3.3-3.4 (3H, m), 3.8 (1H, br s), 5.1 (1H, d), 6.6 (1H, d), 6.9-7.0 (2H, m), 7.05 (1H, d), 7.15-7.22 (5H, m), 7.37 (1H, t); MS: 573 (MH⁺).

Step 2:



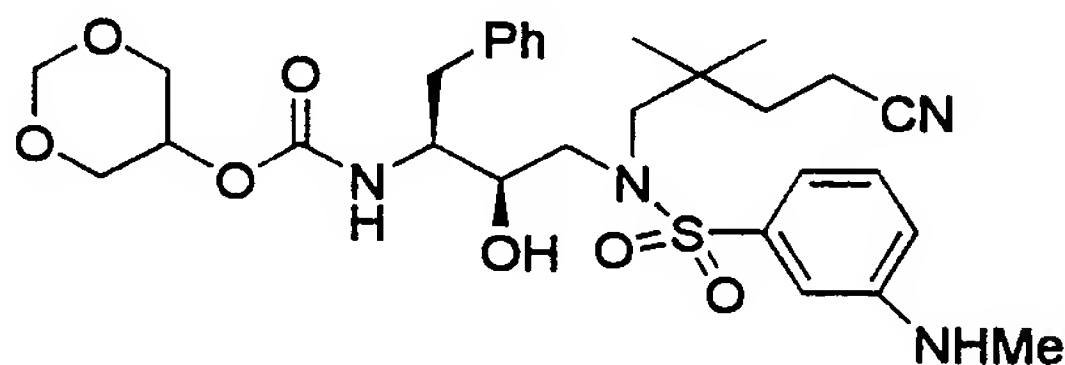
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(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(4-cyano-2,2-dimethylbutyl)[(3-dimethylaminophenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl) carbamate

The product from Step 1 was subjected to
5 procedure similar to Example (Compound 16 to provide the title compound as a foam.

¹H NMR (DMSO-*d*₆): δ 0.9 (3H, s), 0.94 (3H, s), 1.1 (1H, br d), 1.3 (1H, br t), 1.7 (2H, t), 2.4-2.5 (3H, m), 2.6-2.9 (3H, m), 2.95 (6H, s), 3.2-3.4 (2H, m), 3.42-3.45 (1H, m), 3.5-3.6 (2H, m), 3.7 (1H, t), 3.8-3.9 (2H, m), 4.8 (1H, quartet), 5.2 (1H, d), 5.5 (1H, d), 6.9-7.4 (10H, m); MS: 629 (MH⁺).

Example (Compound 24)



1,3-Dioxan-5-yl *N*-[(1*S*,2*R*)-1-benzyl-3-((4-cyano-2,2-dimethylbutyl)[3-(methylamino)phenyl]sulfonylamino)-2-hydroxypropyl] carbamate

The product from step 1 (Example (Compound 12)) was subjected to procedure used in Example (Compound 7) using 1,3-dioxan-5-yl 4-nitrophenylcarbonate to provide
20 the title compound as a foam.

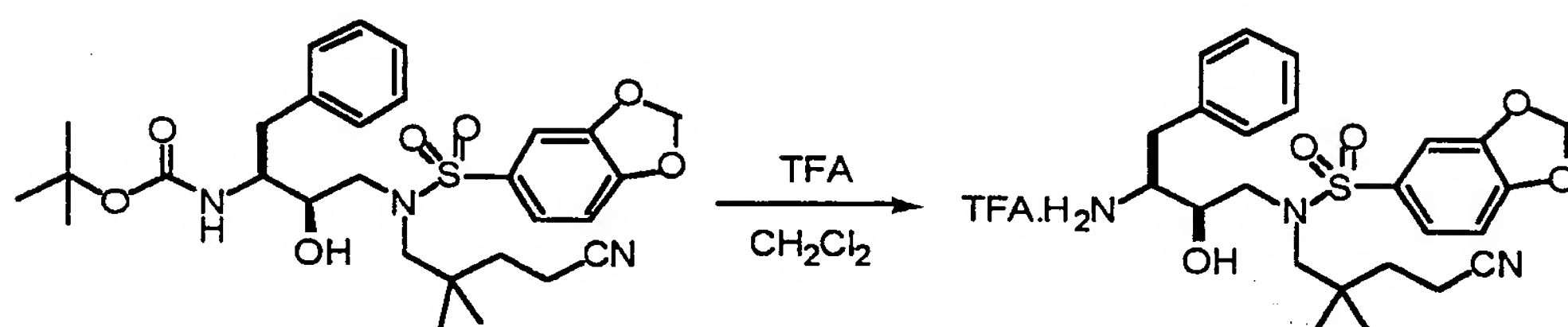
¹H NMR (DMSO-*d*₆): δ 0.9 (3H, s), 0.95 (3H, s), 1.7 (2H, t), 2.4-2.5 (2H, m), 2.7 (3H, d), 2.73 (1H, d), 2.85 (1H, dd), 2.9 (1H, dd), 3.2-3.3 (3H, m), 3.4 (2H, br d), 3.65 (1H dd), 3.77 (1H, dd), 3.8-3.9 (2H, m), 4.25 (1H, br s),
25 4.7 (1H, d), 4.8 (1H, d), 5.1 (1H, d), 6.15 (1H,

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quartet), 6.75 (1H, d), 6.88 (1H, s), 6.92 (1H, d), 7.1-7.3 (7H, m); MS: 589 (MH⁺).

Example (Compound 25)

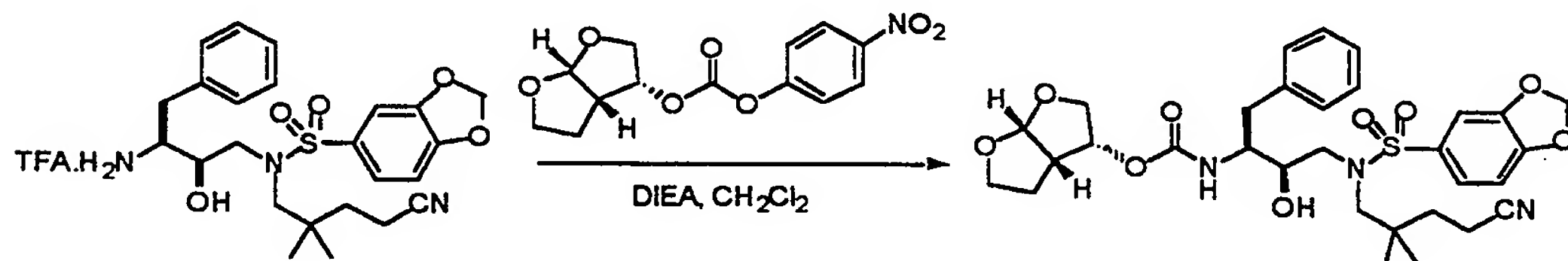
5 Step 1:



10 Benzo[1,3]dioxole-5-sulfonic acid [(2R,3S)-3-amino-2-hydroxy-4-phenyl-butyl]-(4-cyano-2,2-dimethyl-butyl)-amide trifluoroacetate.

A solution of {(1S,2R)-3-[(Benzo[1,3]dioxole-5-sulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid *tert*-butyl ester (0.075 g, 0.13 mmol) in CH₂Cl₂ at ambient temperature was treated
15 with TFA. After stirring 30 min solvent was removed *in vacuo* to give desired product as a white solid which was used directly in the next step.

Step 2:

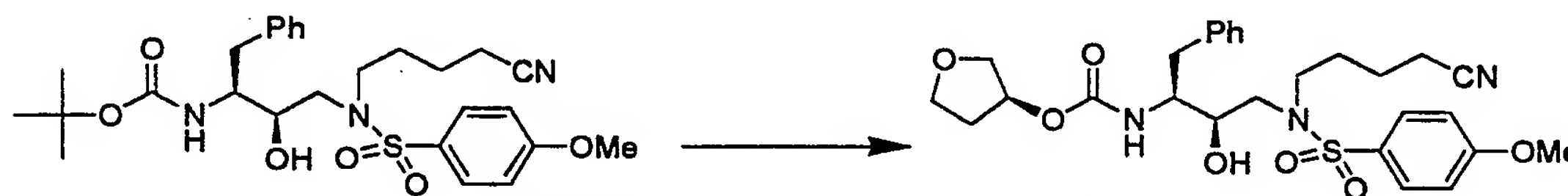


20

{(1S,2R)-3-[(Benzo[1,3]dioxole-5-sulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (3R, 3aS, 6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

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Benzo[1,3]dioxole-5-sulfonic acid [(2R,3S)-3-amino-2-hydroxy-4-phenyl-butyl]-(4-cyano-2,2-dimethyl-butyl)-amide hydrochloride (0.13 mmol) was dissolved in CH₂Cl₂ and treated with diisopropylethylamine (0.114 ml, 0.65 mmol) and (3R, 3aS, 6aR)hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl carbonate (0.046 g, 0.16 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (1% to 2% MeOH in CH₂Cl₂) gave 0.052 g of a white solid. MS (ES): 630 (M+1).

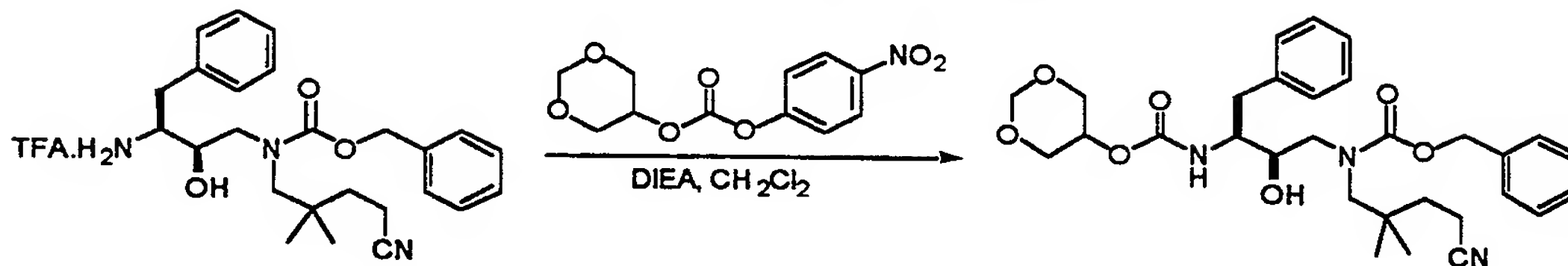
Example (Compound 26)

(3S)-Tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-(4-cyanobutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

The product from Example (Compound 4) was subjected to general procedure listed in Example (Compound 7 to give the title compound as a white glass.

¹H NMR (DMSO-*d*₆): δ 1.4-1.6 (4H, m), 1.7-1.8 (1H, m), 2.0-2.1 (1H, m), 2.4-2.5 (3H, m), 2.75 (1H, dd), 3.0 (2H, br d), 3.2-3.4 (3H, m), 3.5-3.7 (5H, m), 3.8 (3H, s), 4.95 (1H, br s), 5.1 (1H, br s), 7.1 (2H, d), 7.15-7.30 (6H, m), 7.75 (2H, d); MS: 546 (MH⁺).

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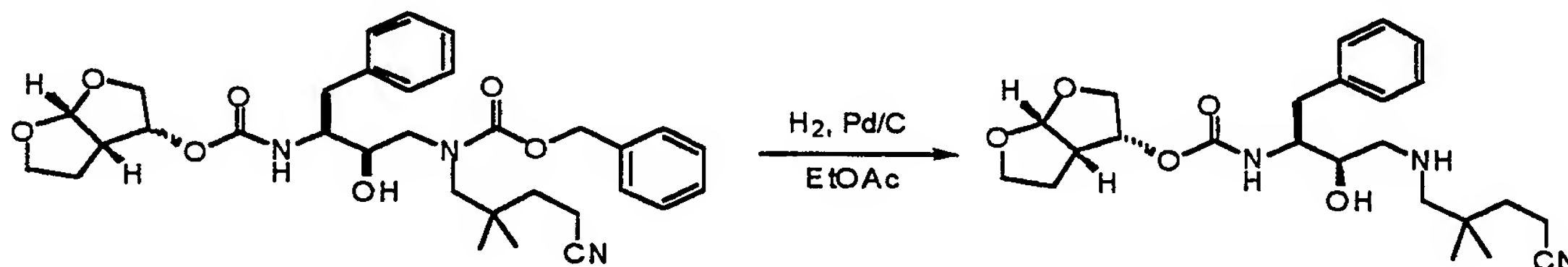
Example (Compound 28)

(4-Cyano-2,2-dimethyl-butyl)-[(2R,3S)-3-([1,3]dioxan-5-ylloxycarbonylamino)-2-hydroxy-4-phenyl-butyl]-carbamic acid benzyl ester.

[(2R,3S)-3-Amino-2-hydroxy-4-phenyl-butyl]-(4-cyano-2,2-dimethyl-butyl)-carbamic acid benzyl ester; hydrochloride (1.30 mmol) was dissolved in CH_2Cl_2 and treated with diisopropylethylamine (0.680 ml, 3.91 mmol) and 1,3-dioxan-5-yl 4-nitrophenyl carbonate (0.421 g, 1.56 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO_3 , and brine. The organic phase was dried over MgSO_4 , filtered and solvent removed *in vacuo*. Purification by column chromatography (3% MeOH in CH_2Cl_2) gave 0.430 g of a white solid. MS (ES): 554 (M+1).

Example (Compound 29)

Step 1:

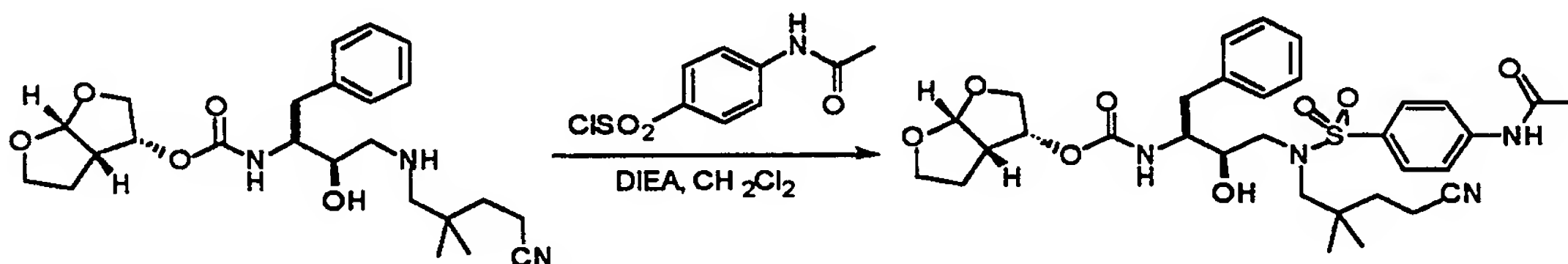


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[(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

A solution of (2S,3R)-(4-Cyano-2,2-dimethyl-butyl)-[3-((3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yloxy-carbonylamino)-2-hydroxy-4-phenyl-butyl]-carbamic acid benzyl ester (0.740 g, 1.28 mmol) in EtOAc at ambient temperature was treated with 10% palladium on carbon (0.075g) and stirred under a balloon of hydrogen. After stirring 16 h the mixture was filtered and solvent was removed *in vacuo* to give desired product as a colorless oil.

Step 2:



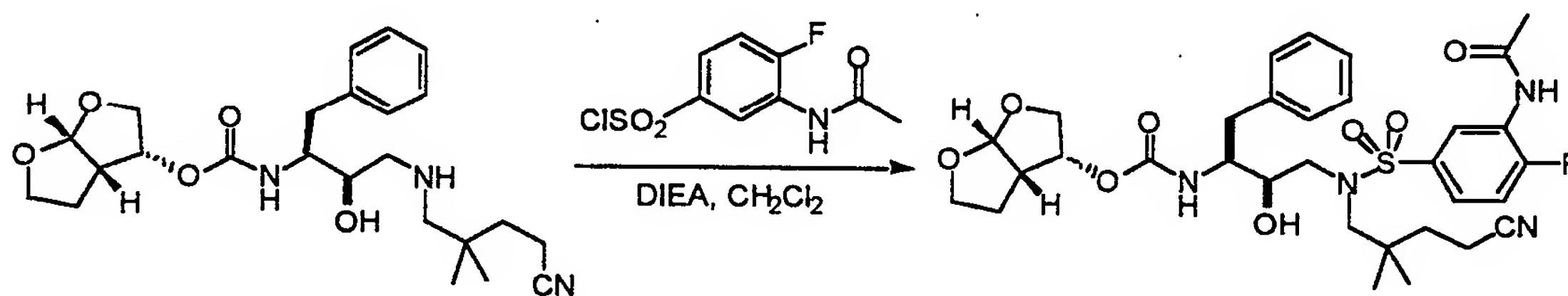
{(1S,2R)-3-[(4-Acetylamino-benzenesulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

[(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester (0.020g, 0.05 mmol) was dissolved in CH₂Cl₂ and treated with diisopropylethylamine (0.016 ml, 0.09 mmol) and 4-acetamidobenzenesulfonyl chloride (0.011 g, 0.05 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated *in vacuo*, taken

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up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by preparative thin layer chromatography (EtOAc) gave 0.006 g of a white solid. MS (ES): 643 (M+1).

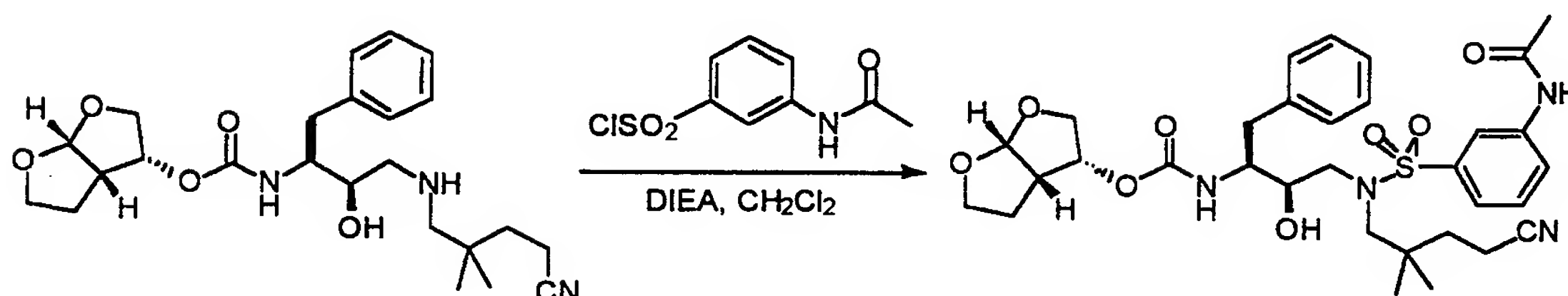
Example (Compound 30)



{ (1S,2R)-3-[(3-Acetylamino-4-fluorobenzenesulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

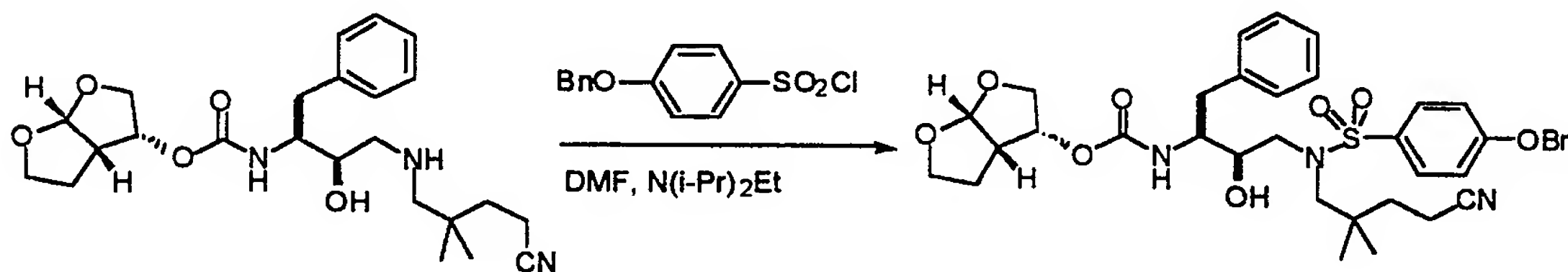
[(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester (0.020g, 0.05 mmol) was dissolved in CH₂Cl₂ and treated with diisopropylethylamine (0.025 ml, 0.13 mmol) and 3-acetamido-4-fluorobenzenesulfonyl chloride (0.012 g, 0.05 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by preparative thin layer chromatography (3% MeOH in CH₂Cl₂) gave 0.013 g of a white solid. MS (ES): 661 (M+1).

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Example (Compound 31)

5 { (1S,2R)-3-[(3-Acetylamino-benzenesulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

[(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester (0.020g, 0.05 mmol) was dissolved in CH₂Cl₂ and treated with diisopropylethylamine (0.025 ml, 0.13 mmol) and 3-acetamidobenzenesulfonyl chloride (0.012 g, 0.05 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by preparative thin layer chromatography (3% MeOH in CH₂Cl₂) gave 0.019 g of a white solid. MS (ES): 643 (M+1).

Example (Compound 32)

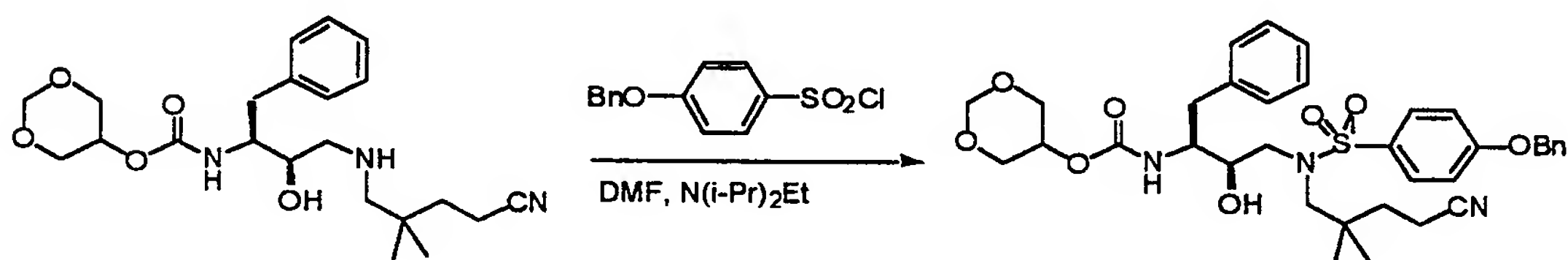
-129-

{ (1S,2R) -3- [(4-benzyloxybenzenesulfonyl) - (4-cyano-2,2-dimethyl-butyl) -amino] -1-benzyl-2-hydroxy-propyl} - carbamic acid (2R, 3aS, 6aR) hexahydro-furo[2,3-b]furan-3-yl ester.

5 [(1S,2R) -1-Benzyl-3- (4-cyano-2,2-dimethyl-butylamino) -2-hydroxy-propyl] -carbamic acid (3R, 3aS, 6aR) hexahydro-furo[2,3-b]furan-3-yl ester (9.5 mg, 0.02 mmol) was combined with 4-benzyloxybenzenesulfonyl chloride (7.2 mg, 1.2 eq.) in
10 anhydrous DMF (1 mL). Solution was chilled to 0 °C and N(*i*-Pr)₂Et (13.4 µL, 3 eq.) was added. Reaction was allowed to warm to room temperature and stirred for 5 hours. Reaction mixture was diluted in EtOAc (10 mL) and washed with sat. NaHCO₃ (10 mL), 0.5N KHSO₄ (10 mL) and
15 brine (10 mL). Organic phase was dried with MgSO₄ and solvent was removed in vacuo. Purification by preparative TLC (1:1, EtOAc:Hex). Recovered 6.1 mg (41%) of the product as a colorless residue. MS (ES): 692.2 (M+1), HPLC; t_R=13.19 min. (94%).

20

Example (Compound 33)



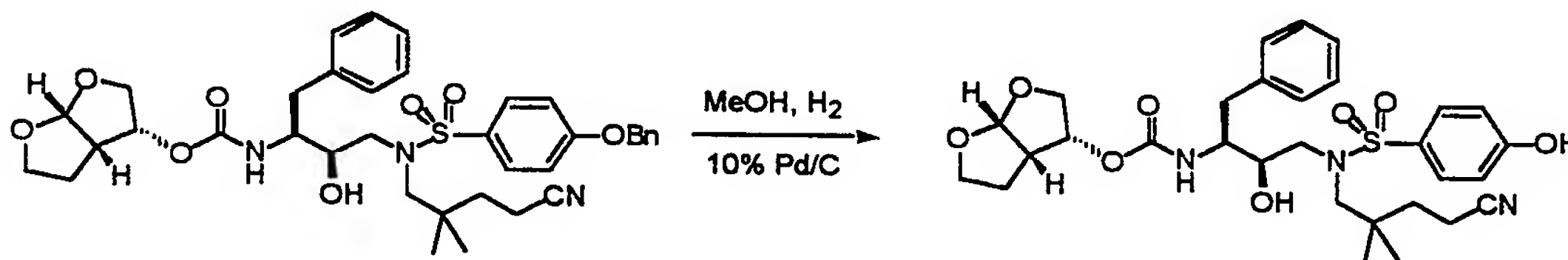
{ (1S,2R) -1-Benzyl-3- [(4-cyano-2,2-dimethyl-butyl) - (4-benzyloxybenzenesulfonyl) -amino] -2-hydroxy-propyl} - carbamic acid [1,3]dioxan-5-yl ester.

25 { (1S,2R) -1-Benzyl-3- (4-cyano-2,2-dimethyl-butylamino) -2-hydroxy-propyl} -carbamic acid [1,3]dioxan-

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5-yl ester (14.8 mg, 0.04 mmol) was combined with 4-benzyloxybenzenesulfonyl chloride (12.0 mg, 1.2 eq.) in anhydrous DMF (1 mL). Solution was chilled to 0 °C and N(*i*-Pr)₂Et (18.4 µL, 3 eq.) was added. Reaction was
 5 allowed to warm to room temperature and stirred for 5 hours. Reaction mixture was diluted in EtOAc (10 mL) and washed with sat. NaHCO₃ (10 mL), 0.5 N KHSO₄ (10 mL) and brine (10 mL). Organic phase was dried with MgSO₄ and solvent was removed in vacuo. Purification by
 10 preparative TLC (1:1, EtOAc:Hex). Recovered 4.2 mg (18%) of the product as a colorless residue. MS (ES); 666.2 (M+1), HPLC; t_R=13.24 min. (98%).

Example (Compound 35)



15 { (1S,2R)-3-[(4-hydroxybenzenesulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (2R, 3aS, 6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

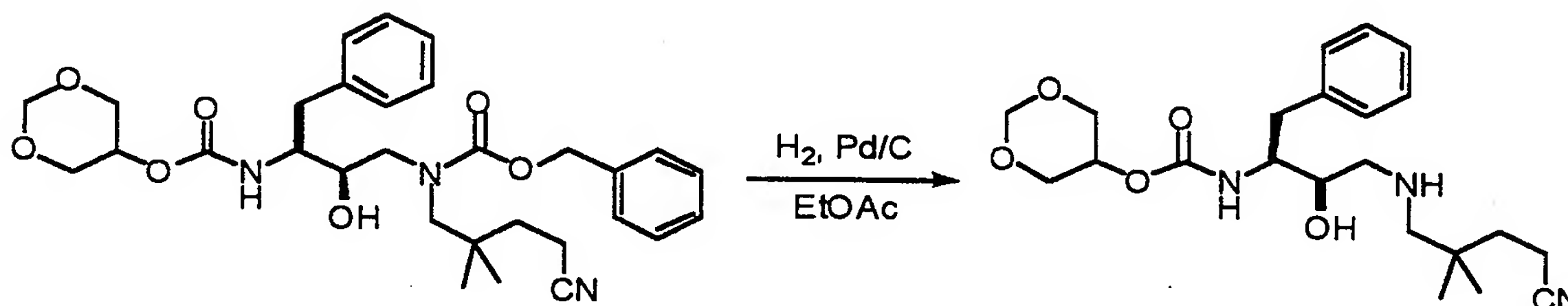
{ (1S,2R)-3-[(4-benzyloxybenzenesulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (2R, 3aS, 6aR)hexahydro-furo[2,3-b]furan-3-yl ester (18.1 mg, 0.03 mmol) was dissolved in methanol (1 mL). Solution was degassed with nitrogen for 5 minutes the 10% Pd/C (2 mg) was added. Reaction was
 20 pressurized with H₂ (1 ATM) and stirred for 30 minutes. Catalyst removed by filtration and solvent was removed in

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vacuo leaving 15.2 mg (97%) of product as a colorless residue. MS (ES): 602 (M+1), HPLC; t_R =10.32 min. (100%).

Example (Compound 36)

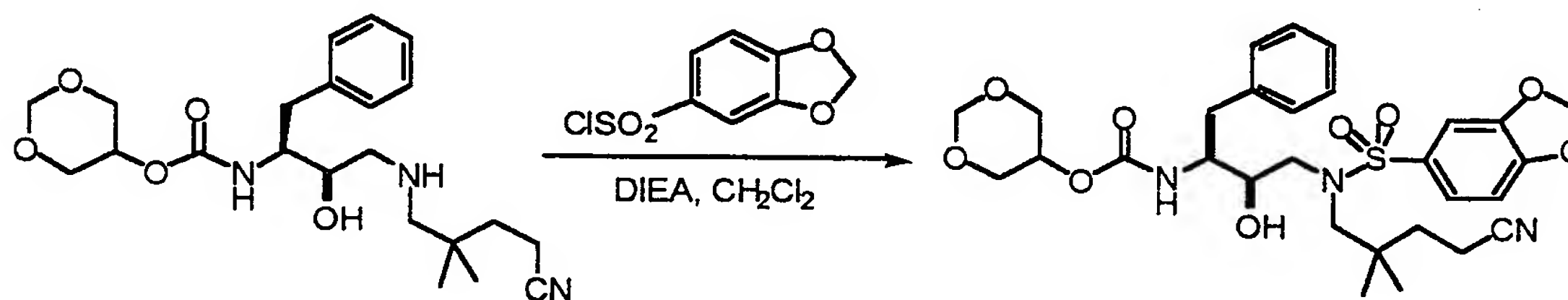
Step 1:



5 [(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-carbamic acid [1,3]dioxan-5-yl ester.

A solution of (4-Cyano-2,2-dimethyl-butyl)-[(2R,3S)-3-([1,3]dioxan-5-yloxycarbonylamino)-2-hydroxy-4-phenyl-butyl]-carbamic acid benzyl ester (0.420 g, 0.76
10 mmol) in EtOAc at ambient temperature was treated with palladium on carbon (0.050g) and stirred under a balloon of hydrogen. After stirring 16 h the mixture was filtered and solvent was removed *in vacuo* to give desired product as a colorless oil.

15 Step 2:



{(1S,2R)-3-[(Benzo[1,3]dioxole-5-sulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid [1,3]dioxan-5-yl ester. [(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-
20 carbamic acid [1,3]dioxan-5-yl ester (0.059g, 0.14 mmol)

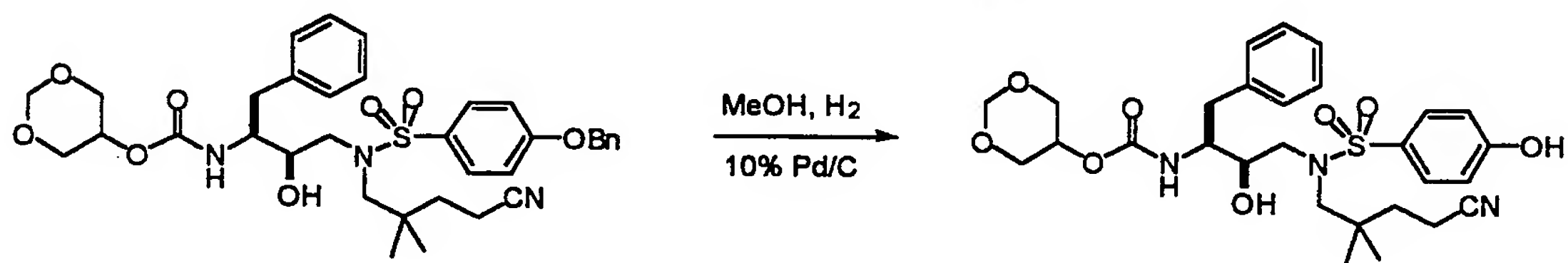
-132-

was dissolved in CH₂Cl₂ and treated with diisopropylethylamine (0.075 ml, 0.42 mmol) and Benzo[1,3]dioxole-5-sulfonyl chloride (0.037 g, 0.17 mmol) at ambient temperature under argon with stirring.

5 After 15 h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (1% to 3% MeOH in CH₂Cl₂) gave

10 0.062 g of a white solid. MS (ES): 604 (M+1).

Example (Compound 42)



{ (1S,2R)-1-Benzyl-3-[(4-cyano-2,2-dimethyl-butyl)-(4-hydroxybenzenesulfonyl)-amino]-2-hydroxy-propyl}-carbamic acid [1,3]dioxan-5-yl ester.

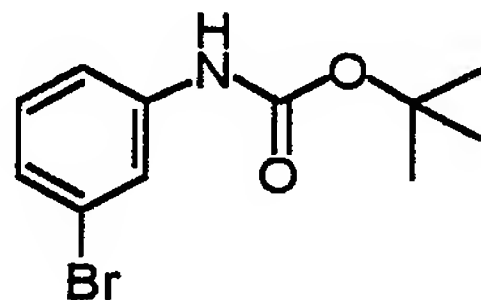
15 { (1S,2R)-1-Benzyl-3-[(4-cyano-2,2-dimethyl-butyl)-(4-benzyloxybenzenesulfonyl)-amino]-2-hydroxy-propyl}-carbamic acid [1,3]dioxan-5-yl ester (2.2 mg, 0.003 mmol) was dissolved in methanol (1 mL). Solution was degassed with nitrogen for 5 minutes the 10% Pd/C (2

20 mg) was added. Reaction was pressurized with H₂ (1 ATM) and stirred for 30 minutes. Catalyst removed by filtration and solvent was removed *in vacuo* leaving 2.0 mg (95%) of product as a colorless residue. MS (ES): 576.2 (M+1), HPLC; *t_R*=10.39 min. (84%).

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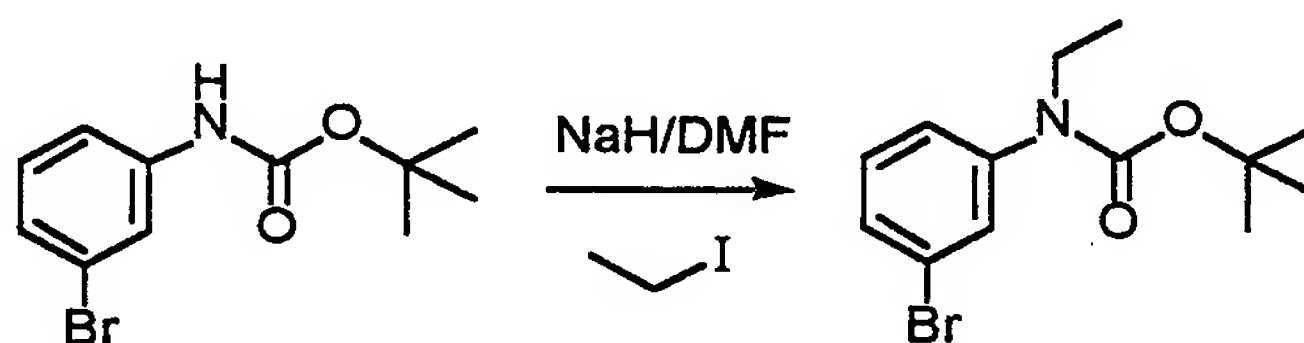
Example (Compound 43)

Step 1:

**3-Bromo-N-tert-butoxycarbonylaniline.**

5 3-Bromoaniline (0.50 ml, 4.6 mmol) di-tert-butylidicarbonate (1.20 g, 5.5 mmol) and 4-dimethylaminopyridine (0.003 g) were combined in anhydrous CH₂Cl₂ (10 ml). Solution chilled to 0°C and triethylamine (1.28 ml, 9.2 mmol) was added. Reaction
10 was allowed to warm to room temperature then was heated to reflux for 1 hour. Reaction mixture was diluted in EtOAc and washed with sat. NaHCO₃, 0.5 N KHSO₄ and brine. Organic phase was dried with MgSO₄ and the solvent was removed *in vacuo*. Purification by flash chromatography
15 (1:4, EtOAc: Hex., gradient to 1:3 then 1:2). Recovered 1.01 g (81%) of the product as a light yellow solid. R_f=0.62 (1:4, EtOAc:Hex). ¹H NMR (CDCl₃) 7.68 (1H, s), 7.22-7.10 (3H, m), 6.48 (1H, b), 1.51 (9H, s).

Step 2:



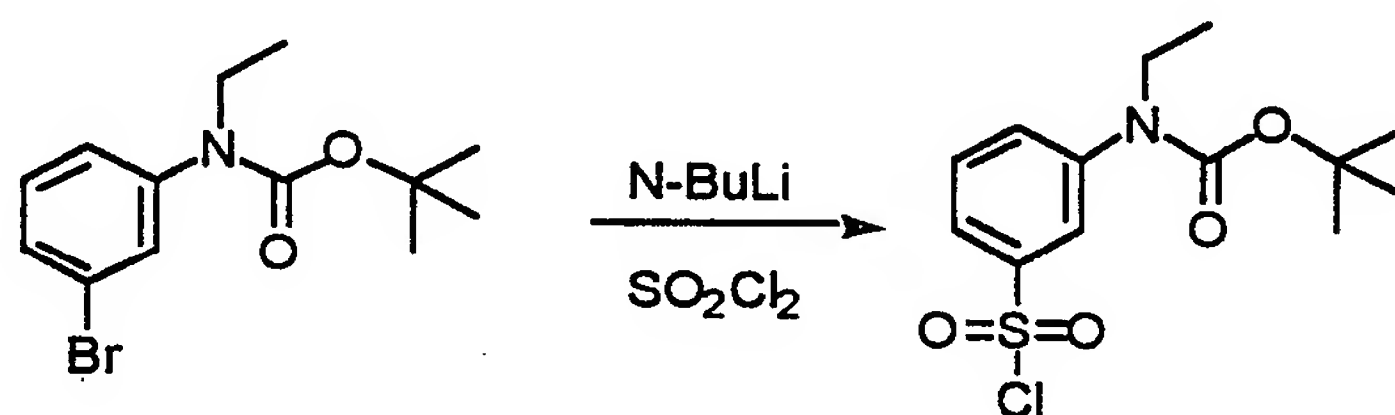
20

3-Bromo-N-tert-butoxycarbonyl-N-ethylaniline.

3-Bromo-N-tert-butoxycarbonylaniline (0.82 g, 3.0 mmol) was dissolved in anhydrous DMF (15 ml) and

-134-

cooled to 0 °C. Sodium hydride (0.145 g, 3.6 mmol, 1.2 eq.) was added to the solution. The reaction stirred at 0 °C for 10 minutes then was warmed to room temperature. After 30 minutes at room temperature the reaction was
5 cooled to 0 °C and Ethyl iodide (0.27 ml, 3.3 mmol, 1.1 eq.) was added over ~ 1 minute. The reaction was stirred at 0 °C for 5 minutes then was warmed to room temperature. The reaction stirred at room temperature for 22 hours and was then cooled to 0 °C and quenched with a saturated
10 bicarbonate solution and extracted with EtOAc (2 times). The combined organic layers were washed with H₂O (5 times) then brine. The organic phase was dried with MgSO₄/Na₂SO₄, filtered and the solvent was removed in vacuo to give 0.93 g of crude product which
15 was used as is. HPLC ret time = 11.83 min. (100%).

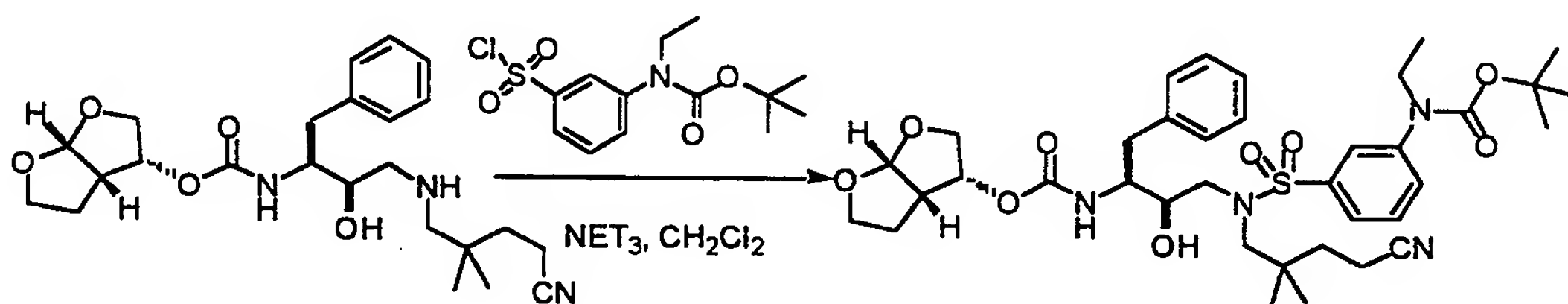
Step 3:**N-tert-Butoxycarbonyl-3-chlorosulfonyl-N-ethylaniline.**

3-Bromo-N-tert-butoxycarbonyl-N-ethylaniline
20 (3.0 mmol) was dissolved in freshly distilled THF (25 ml) under a N₂ atmosphere. The solution was chilled to -78 °C and butyl lithium (1.8 ml, 2.0 M solution in cyclohexane, 3.6 mmol) was added. After 35 minutes, sulfuryl chloride (0.30 ml, 3.6 mmol) was added over ~ 2 minutes. The
25 reaction was stirred for 5 minutes at -78 °C then was

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allowed to warm to room temperature. After 30 minutes at room temperature the reaction was cooled to -78°C and quenched with a saturated ammonium chloride solution. The THF was removed by evaporation and the resulting residue was and partitioned between EtOAc and H₂O. The aqueous phase was extracted with EtOAc and the combined organic phases were washed with H₂O and brine before being dried with Na₂SO₄. The solvent was removed *in vacuo* to give 900 mg of crude product. HPLC showed the material to be ~ 80% pure and it was used as is. LC/MS (ES): 320.1 (M+1).

Step 4:



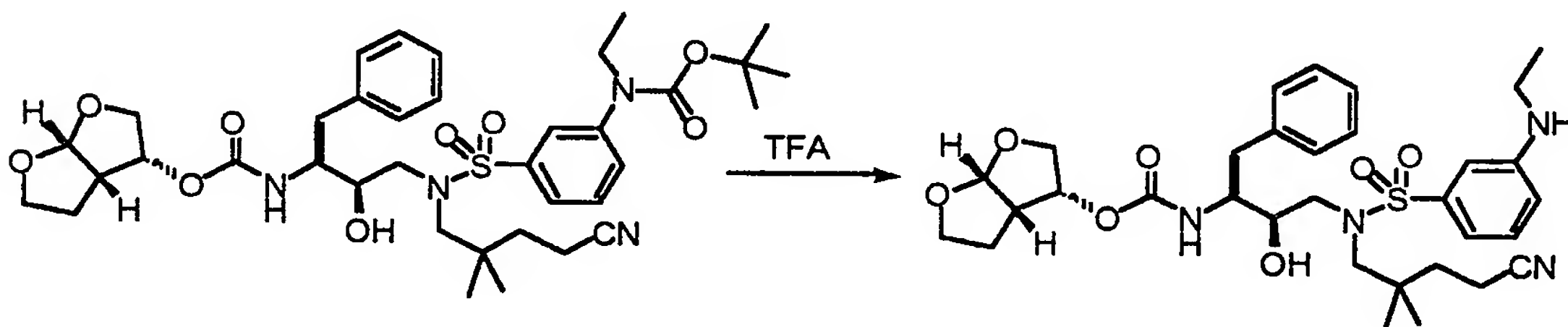
{ (1S,2R)-3-[(3-N-tert-butoxycarbonyl-N-ethylaminophenyl)sulfonyl]-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid
(3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

[(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-carbamic acid
(3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester
(0.025g, 0.056 mmol) was dissolved in 1 mL of CH₂Cl₂ and treated with triethylamine (0.032 ml, 0.22 mmol). A solution N-tert-Butoxycarbonyl-3-chlorosulfonyl-N-ethylaniline (0.045 g, 0.14 mmol) in 1 mL of CH₂Cl₂ was added followed by 4-dimethylaminopyridine (0.005 g). The reaction was stirred at room temperature under N₂ for 23

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hours, then quenched with a saturated bicarbonate solution and extracted with EtOAc (2 times). The combined organic layers were washed with H₂O, then brine. The organic phase was dried over MgSO₄, filtered and solvent removed in vacuo. Purification by preparative thin layer chromatography (70:30:2, EtOAc:Hexanes:MeOH) gave 0.0018 g of product. HPLC ret time = 11.49 min. LC/MS (ES); = 729.3 (M+1).

Example (Compound 44)



{ (1S,2R)-3-[(3-N-ethylaminophenyl)sulfonyl]-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl} carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

{ (1S,2R)-3-[(3-N-tert-butoxycarbonyl-N-ethylaminophenyl)sulfonyl]-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester (0.007 g, 0.009 mmol) was dissolved in 0.5 mL of CH₂Cl₂. Trifluoroacetic acid (0.25 mL) was added and the reaction was stirred for 1 hour at room temperature under N₂. The solvent was removed in vacuo and the residue was partitioned between EtOAc and a saturated bicarbonate solution. The aqueous phase was extracted with EtOAc and the combined organic phases were washed with H₂O and brine before being dried with Na₂SO₄ and filtered. The solvent

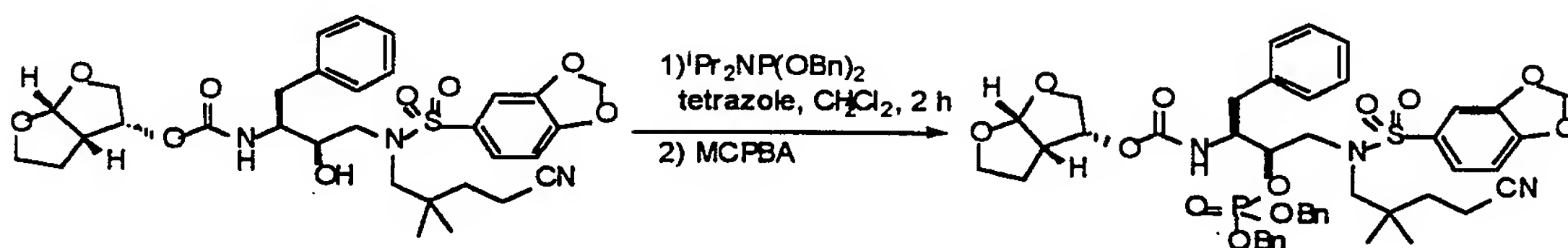
-137-

was removed *in vacuo* to give the 2.1 mg of product.

HPLC ret. time = 9.156 min. LC/MS (ES) = 629.3 (M+1).

Example (Compound 47)

5 Step 1:



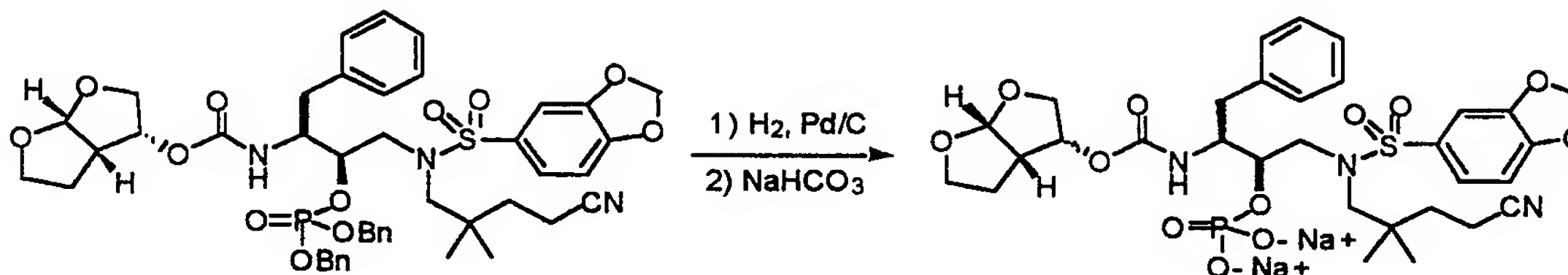
[(1S,2R)-3-[(Benzo[1,3]dioxole-5-sulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-(dibenzylphosphoryloxy)-propyl]-carbamic acid

10 (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

A solution of {(1S,2R)-3-[(Benzo[1,3]dioxole-5-sulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid (3R, 3aS, 6aR)hexahydro-furo[2,3-b]furan-3-yl ester (0.077 g, 0.12 mmol) and 1H-tetrazole (0.012g, 0.17 mmol) in CH₂Cl₂ was treated with dibenzyl diisopropylphosphoramidite (0.050 ml, 0.015 mmol) and stirred at ambient temperature under argon. After 2 h 3-chloroperoxybenzoic acid (0.072g, 0.30 mmol) was added and the reaction mixture stirred an additional 1 h. Solvent was removed *in vacuo*, the residue taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo* to give desired product, which was used directly in the next step.

25 Step 2:

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5 { (1S,2R)-3-[(Benzo[1,3]dioxole-5-sulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-phosphonooxy-propyl}-carbamic acid (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester; disodium salt.

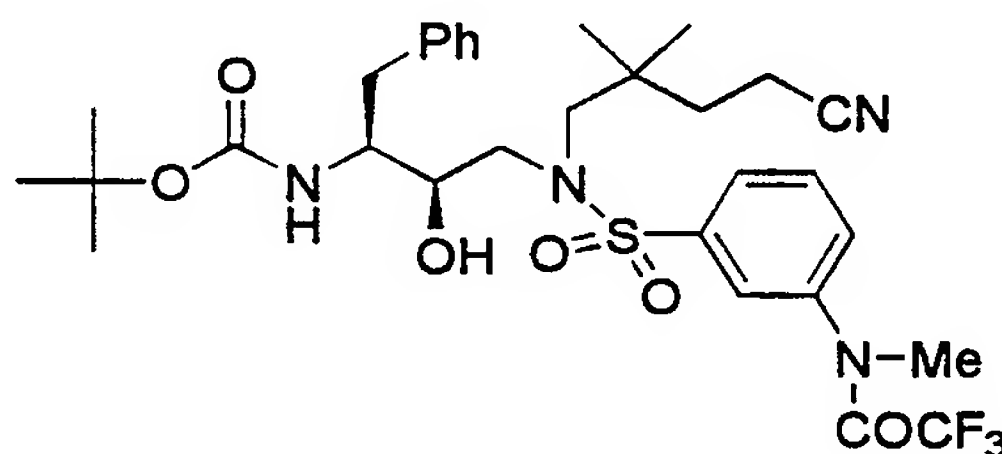
[(1S,2R)-3-[(Benzo[1,3]dioxole-5-sulfonyl)-(4-cyano-2,2-dimethyl-butyl)-amino]-1-benzyl-2-(dibenzoyloxy-phosphoryloxy)-propyl]-carbamic acid

(3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester (0.081 g, 0.09 mmol) was dissolved in EtOAc and treated with 10% palladium on carbon (0.020g) and stirred under a balloon of hydrogen. After stirring 14 h the mixture was filtered and solvent was removed *in vacuo*. The crude residue was purified by preparative HPLC (15% to 80% CH₃CN in H₂O) and the pooled fractions concentrated *in vacuo* to give 0.029g of product. This was taken up in 0.10 M sodium bicarbonate (2.0 equiv., 0.817 ml, 0.09 mmol) stirred 2 h at ambient temperature and lyophilized to give 0.029 g of the bis sodium salt as a white solid.

20

Example (Compound 100)

Step 1:



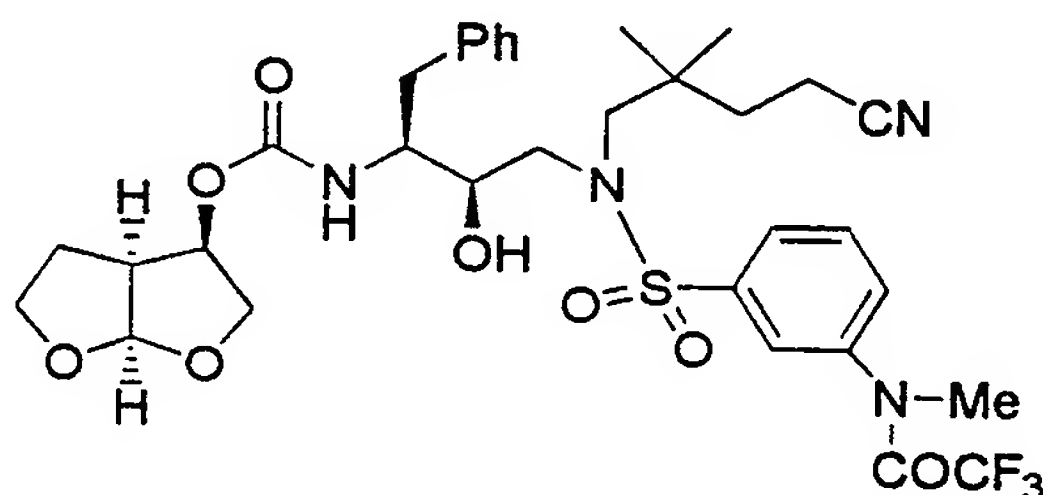
-139-

ter-Butyl N-((1*S*,2*R*)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)[(3-(*N*-methyl,*N*-trifluoroacetyl)aminophenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

To a solution of the product from Example
5 (Compound 6), 0.44 g, 0.8 mmol), in dichloromethane (17 mL), and pyridine (1 mL, 0.97 g, 12.3 mmol) at 0°C was added trifluoroacetic anhydride (0.45 mL, 0.67 g, 3.2 mmol) and the mixture was stirred at ambient temperature for 2 hours. The mixture was washed with saturated
10 sodium bicarbonate, dried (sodium sulfate) and coevaporated with ethyl acetate (3X). The residue was dissolved in methanol (10 mL) and potassium carbonate (10 mg) was added. After stirring for 1 hour at ambient temperature, the solvent was evaporated and the residue
15 was dissolved in acetone (15 mL) and potassium carbonate (0.26 g, 2 mmol) and methyl iodide (0.25 mL, 0.56 g, 4.0 mmol) were added. After 3 hours, the solvent was evaporated and the residue was partitioned between dichloromethane and water. The organic phase was dried
20 (sodium sulfate), evaporated, dried in vacuo to provide the title compound (0.52 g) as a foam.

¹H NMR (CDCl₃): δ 1.0 (6H, s), 1.14 (9H, s), 1.75 (2H, t), 2.38 (2H, t), 2.7-2.8 (1H, m), 2.9 (1H, dd), 3.05-3.25 (3H, m), 3.4 (3H, s), 3.5-3.7 (3H, m), 3.9 (1H, br s),
25 4.5 (1H, br s), 7.1-7.3 (5H, m), 7.5 (1H, d), 7.6 (1H, t), 7.7-7.9 (2H, m); MS: 677 (M + 23)

Step 2:



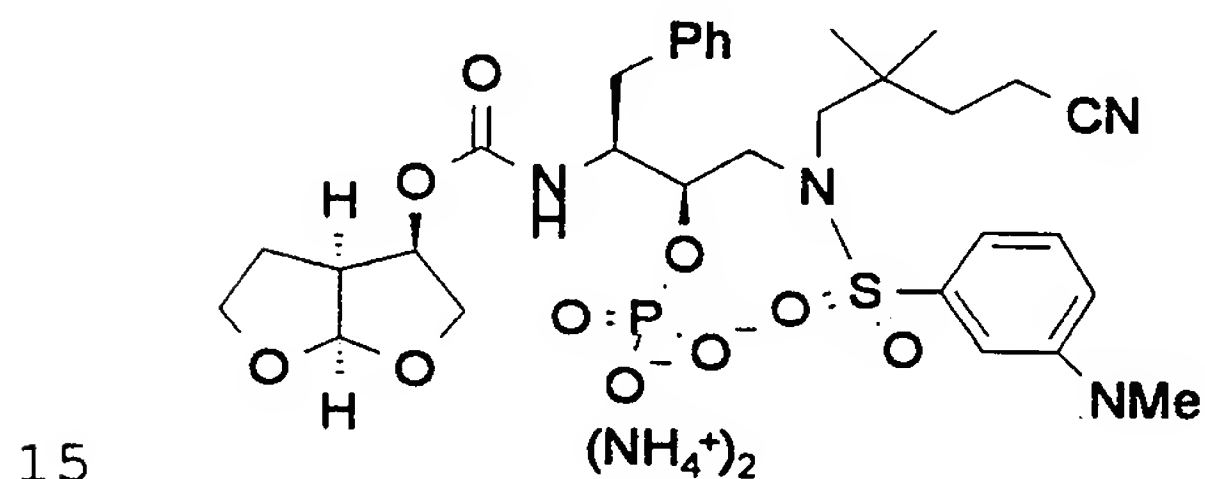
-140-

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-benzyl-3-((4-cyano-2,2-dimethylbutyl)[3-(N-methyl,N-trifluoroacetylmethylamino)phenyl]sulfonylamino)-2-hydroxypropyl]carbamate

5 The product from step 1 was subjected to the procedure outlined in Example (Compound 10) to provide the title compound as a foam.

¹H NMR (DMSO-d₆): δ 0.93 (3H, s), 0.96 (3H, s), 1.14 (1H, dd), 1.33-1.36 (1H, m), 1.65 (2H, t), 2.4-2.5 (3H, m), 2.6-2.8 (2H, m), 2.9-3.1 (2H, m), 3.3 (3H, s), 3.32-3.60 (5H, m), 3.65-3.82 (3H, m), 4.8 (1H, quartet), 5.1 (1H, d), 5.5 (1H, d), 7.1-7.3 (6H, m), 7.7 (1H, t), 7.79 (1H, d), 7.88 (1H, d), 8.0 (1H, s); MS: 711 (MH⁺).

Step 3:



(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)[(3-methylaminophenyl)sulfonyl]amino-2-hydroxypropylcarbamate bis-ammonium phosphate

20 The product from step 2 was subjected to procedures listed in Example (Compound 101) to provide the title compound as a tan solid.

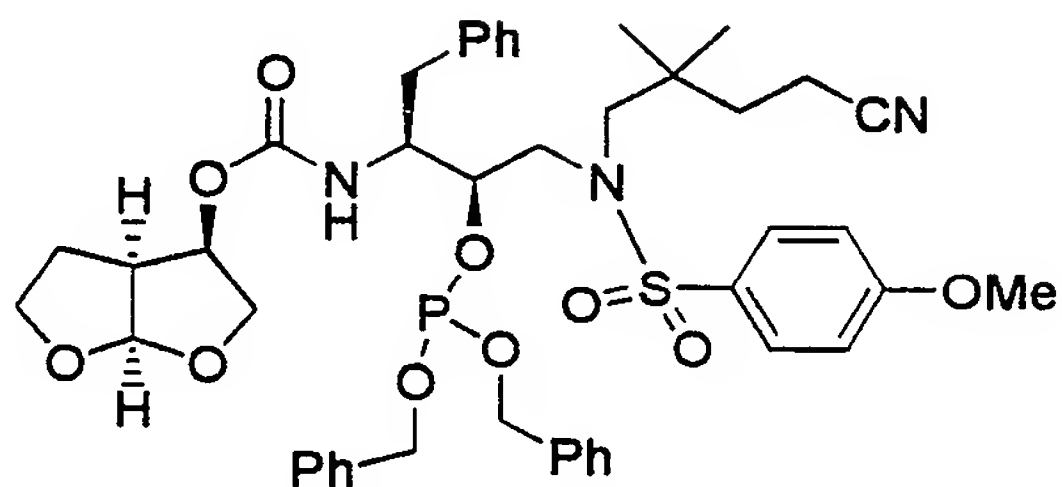
¹H NMR (DMSO-d₆): δ 0.93 (3H, s), 0.96 (3H, s), 1.15-1.40 (2H, m), 1.6 (2H, t), 2.43 (2H, t), 2.6-3.0 (8H, m), 3.2-

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3.6 (4H, m), 3.7 (1H, t), 3.8 (1H, t), 4.0 (1H, br s), 4.35 (1H, br s), 4.4-4.5 (1H, m), 4.8 (1H, quartet), 5.5 (1H, d), 6.7 (1H, d), 6.9 (1H, d), 7.0 (1H, s), 7.1-7.4 (7H, m); MS: 695 (MH⁺);

5 Example (Compound 101)

Step 1:



(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl) [(4-methoxyphenyl) sulfonyl]-2-dibenzylphosphite-aminopropyl) carbamate

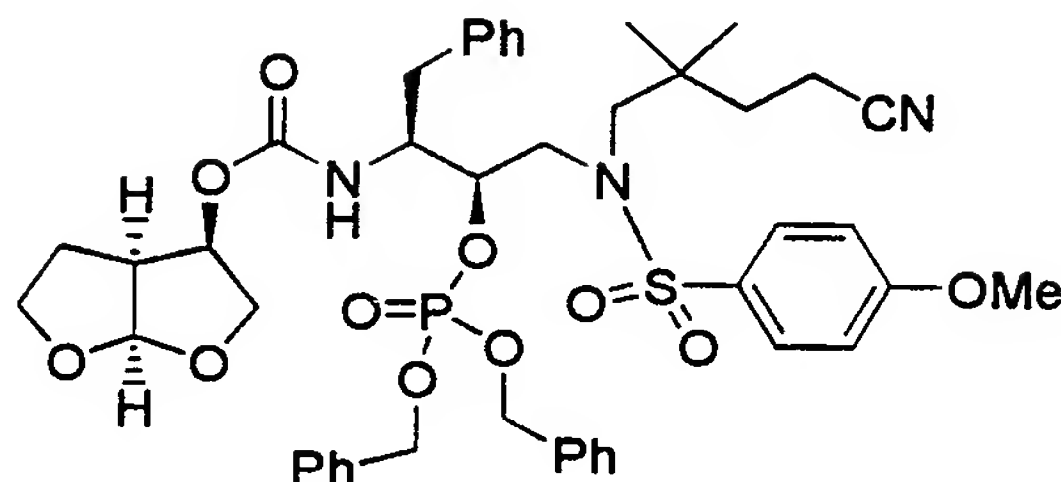
10 To a solution of the product from Example (Compound 21), (0.22 g, 0.36 mmol) in dichloromethane (5 mL) was added tetrazole (32 mg, 0.47 mmol) and dibenzyl diisopropyl phosphoramidite (0.151 mL, 0.155 mg, 0.45 mmol) and the mixture was stirred at ambient temperature
15 under nitrogen atmosphere for 18 hours. The mixture was washed with saturated sodium bicarbonate/water, dried (sodium sulfate) and evaporated. The residue was chromatographed (silica gel, hexane/acetone, 2:1) to provide the title compound as a solid (0.23 g);

20 ¹H NMR (DMSO-d₆): δ 0.8 (3H, s), 0.86 (3H, s), 1.33 (1H, dd), 1.28-1.40 (1H, m), 1.55-1.70 (2H, m), 2.35 (2H, td), 2.4-2.5 (1H, m), 2.73 (1H, br quartet), 2.85 (1H, dd), 3.05 (1H, dd), 3.2-3.3 (3H, m), 3.5 (1H, dd), 3.55 (1H, dd), 3.66 (1H, dd), 3.74 (1H, dd), 3.8 (3H, s), 3.84-3.94

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(1H, m), 4.6-4.7 (1H, m), 4.75 (1H, quartet), 4.8-4.9 (4H, m), 7.1-7.2 (8H, m), 7.23-7.33 (10H, m), 7.8 (2H, d); MS: 882 (M + 23).

Step 2:



- 5 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]-2-dibenzylphosphate-aminopropyl) carbamate

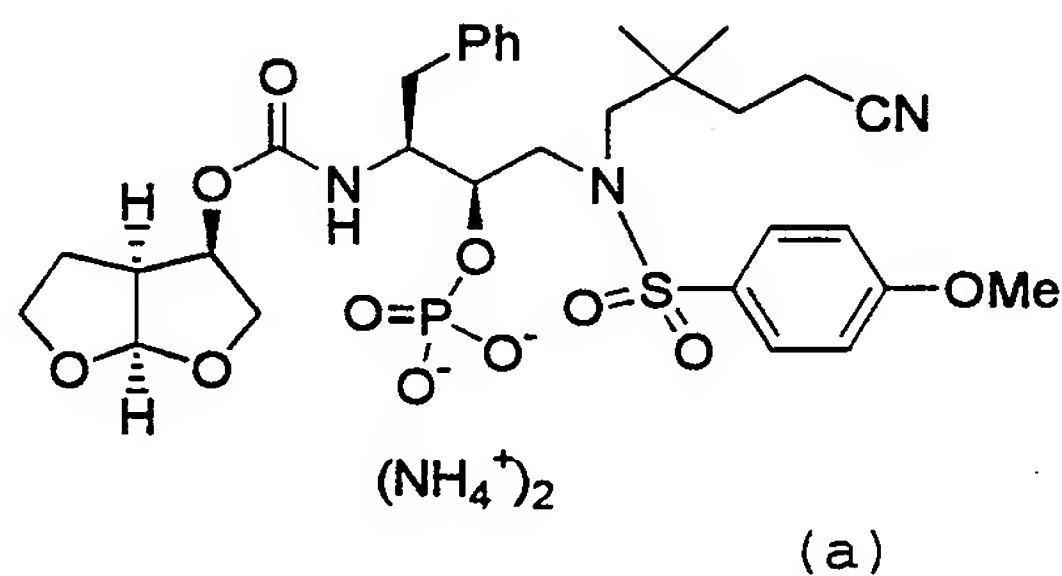
To a solution of the product from step 1 (0.18 g, 0.21 mmol) in anhydrous acetonitrile (5 mL) was added iodobenzene diacetate (0.10 g, 0.31 mmol) and the mixture was stirred at ambient temperature for 1 hour. Solvent was evaporated and the residue was dissolved in dichloromethane, washed with saturated sodium bicarbonate/water, dried (sodium sulfate) and evaporated. The residue was chromatographed (silica gel, dichloromethane/methanol, 98.5:1.5) to give the title compound as a foam (85 mg);

¹H NMR (CDCl₃): δ 0.83 (3H, s), 0.89 (3H, s), 1.38 (1H, dd), 1.5-1.7 (3H, m), 2.2 (2H, t), 2.54 (1H, d), 2.66 (1H, dd), 2.82-2.94 (2H, m), 3.0 (1H, m), 3.22 (1H, d), 3.38 (1H, dt), 3.6-3.7 (2H, m), 3.8 (1H, td), 3.83 (3H, s), 3.9 (1H, dd), 4.15 (1H, td), 4.97 (1H, quartet), 5.0-

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5.2 (4H, m), 5.62 (1H, d), 6.0 (1H, d), 6.98 (2H, d),
7.10-7.25 (6H, m), 7.4 (10H, s), 7.7 (1H, d).

Step 3:

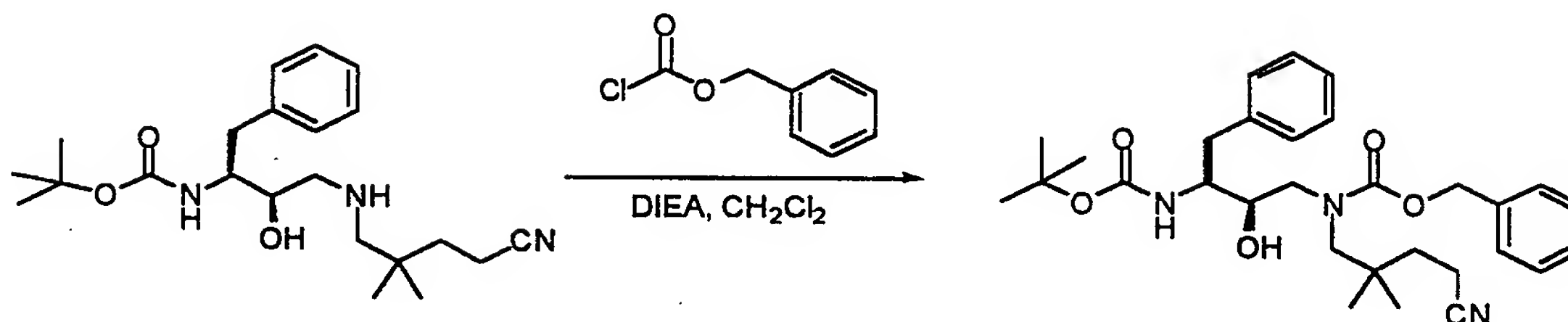


(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(4-cyano-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate-bis-ammonium phosphate

To a solution of the product from step 2 (0.08 g) in 2M ammonia/methanol (5 mL) was added 10 Pd/C (0.02 g) and the mixture was hydrogenated at atmospheric pressure for 4 hours. The mixture was filtered through a bed of celite. Solvent was evaporated and the residue was dried in vacuo to provide the title compound as a solid.

¹H NMR (DMSO-d₆): δ 0.92 (3H, s), 0.95 (3H, s), 1.2-1.4 (2H, m), 1.6 (2H, t), 2.4-2.5 (3H, m), 2.6 (1H, dd), 2.7-2.9 (3H, m), 3.3-3.4 (3H, m), 3.5 (1H, dd), 3.55 (1H, dd), 3.7 (1H, td), 3.8 (3H, s), 3.9 (1H, br t), 4.2 (1H, br s), 4.8 (1H, quartet), 5.5 (1H, d), 7.05 (2H, d), 7.10-7.22 (6H, m), 7.75 (2H, d); MS: 694 (MH⁺).

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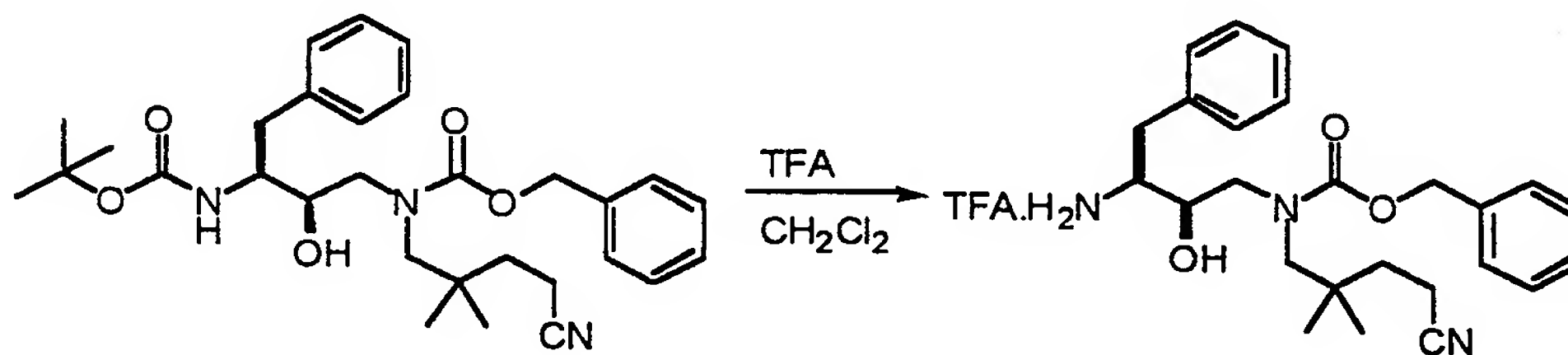
Example (Compound 102)

5 **[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-butyl]-(4-cyano-2,2-dimethyl-butyl)-carbamic acid benzyl ester.**

[(1S,2R)-1-Benzyl-3-(4-cyano-2,2-dimethyl-butylamino)-2-hydroxy-propyl]-carbamic acid *tert*-butyl ester (1.300 g, 3.34 mmol) was dissolved in CH₂Cl₂ and treated with diisopropylethylamine (1.740 ml, 10.01 mmol) and benzyl chloroformate (0.683 g, 4.00 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (10% Et₂O in CH₂Cl₂) gave 0.083 g of a white solid. MS (ES): 524 (M+1).

Example (Compound 103)

Step 1:

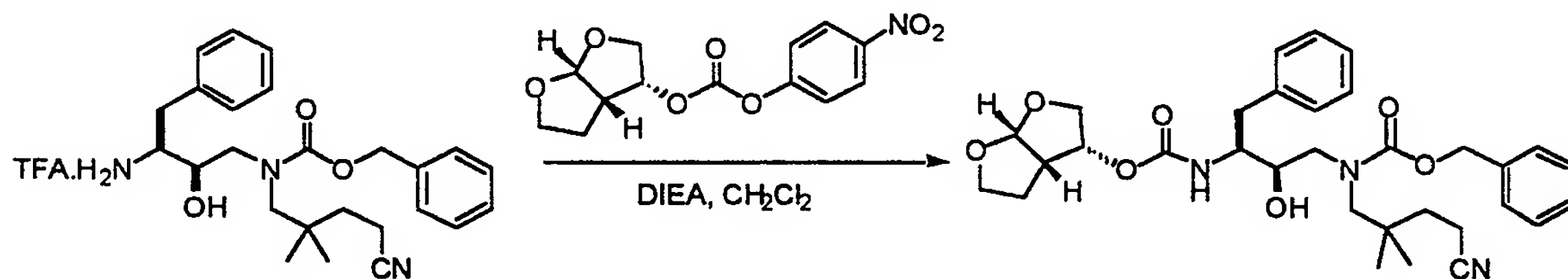


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[(2R,3S)-3-Amino-2-hydroxy-4-phenyl-butyl]-(4-cyano-2,2-dimethyl-butyl)-carbamic acid benzyl ester; trifluoroacetate.

A solution of [(2R,3S)-3-tert-butoxycarbonylamino-2-hydroxy-4-phenyl-butyl]-(4-cyano-2,2-dimethyl-butyl)-carbamic acid benzyl ester (1.291 g, 2.47 mmol) in CH₂Cl₂ at ambient temperature was treated with TFA. After stirring 30 min solvent was removed in vacuo to give desired product as a white solid which was used directly in the next step.

Step 2:



(2S,3R)-(4-Cyano-2,2-dimethyl-butyl)-[3-((3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl)oxycarbonylamino]-2-hydroxy-4-phenyl-butyl]-carbamic acid benzyl ester.

[(2R,3S)-3-Amino-2-hydroxy-4-phenyl-butyl]-(4-cyano-2,2-dimethyl-butyl)-carbamic acid benzyl ester; hydrochloride (2.47 mmol) was dissolved in CH₂Cl₂ and treated with diisopropylethylamine (1.300 ml, 7.40 mmol) and (3R, 3aS, 6aR)hexahydrofuro[2,3-b]furan-2-yl 4-nitrophenyl carbonate (0.801 g, 2.71 mmol) at ambient temperature under argon with stirring. After 15 h the reaction mixture was concentrated in vacuo, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent

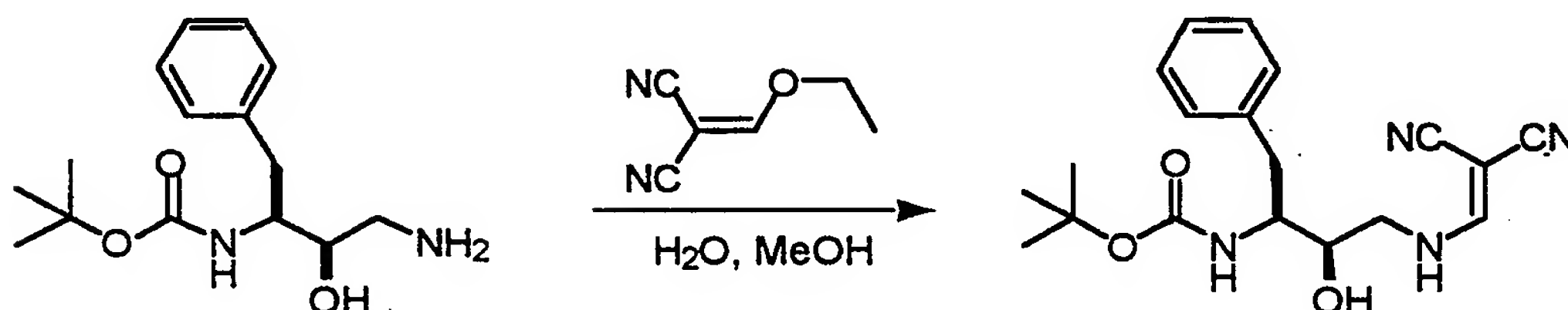
-146-

removed *in vacuo*. Purification by column chromatography (3% MeOH in CH₂Cl₂) gave 0.747 g of a white solid.

MS (ES): 580 (M+1).

Example (Compound 113)

5 Step 1:

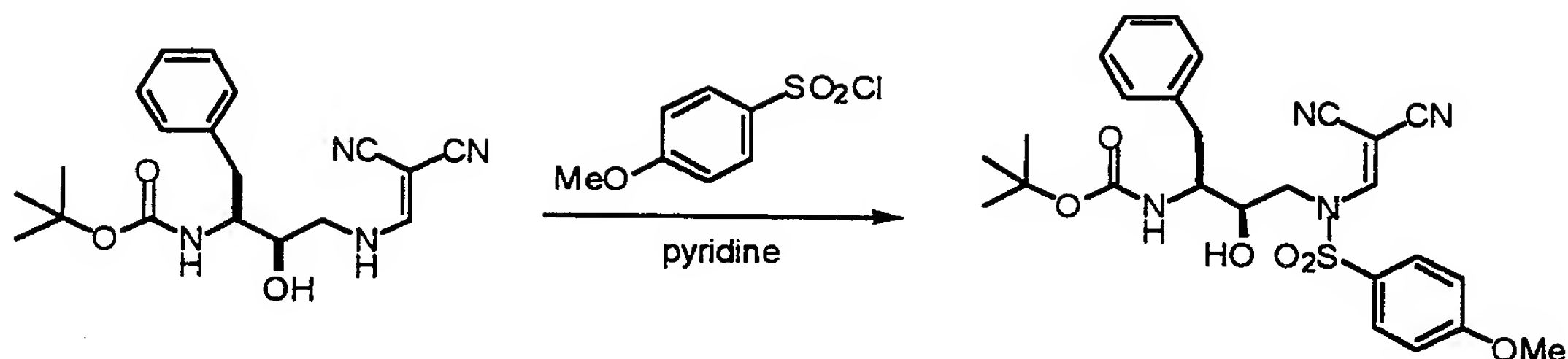


tert-Butyl-N-[(1S,2R)-1-benzyl-3-amino methylene malonitrile-2-hydroxypropyl]carbamate.

tert-Butyl-N-[(1S,2R)-1-benzyl-3-amino-2-hydroxy]carbamate (0.87 mmol) was dissolved in 10 mL of MeOH containing 1% of water and ethoxymethylene malonitrile (107 mg, 0.87 mmol) at ambient temperature with stirring. After 1 hr the reaction mixture was dissolved in EtOAc, washed with water and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (EtOAc 50% in Hexane) gave 0.282 g as an oil. MS (ES): 357.3 (M+1), HPLC showed the material to be 97% pure; ret. time = 11.6 min.

20 Step 2:

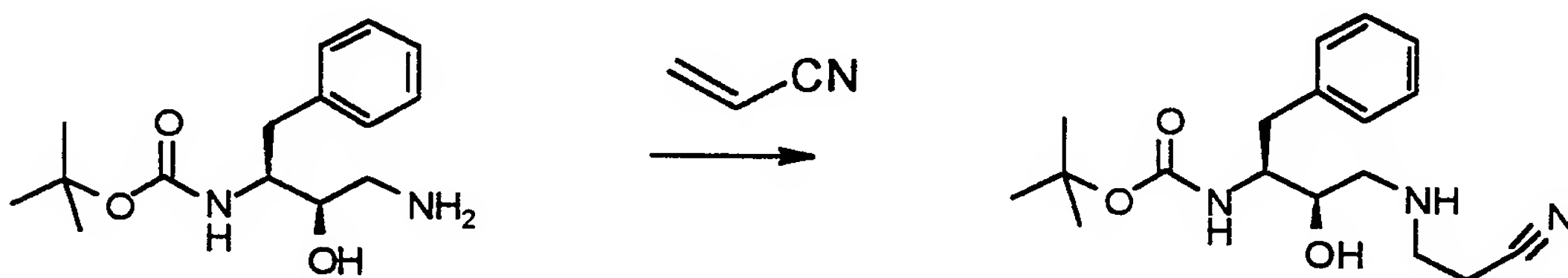
-147-



tert-Butyl-N-[(1*S*,2*R*)-1-benzyl-3-(methylene malonitrile)-2-hydroxypropyl] [(4-methoxyphenyl) sulfonyl]amino] carbamate.

- 5 *tert*-Butyl-N-[(1*S*,2*R*)-1-benzyl-3-amino methylene malonitrile-2-hydroxypropyl]carbamate (0.36 mmol) was dissolved in 2 mL of pyridine. *p*-Methoxybenzene sulfonylchloride (0.09 g, 0.43 mmol) was added and the reaction was stirred at ambient temperature for 3 days.
- 10 The reaction mixture was dissolved in dichloromethane, washed with HCl (1*N*) and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by column chromatography (EtOAc, 50% in Hexane) gave 0.017 g as a solid. MS (ES): 527.3 (*M*+1),
- 15 HPLC showed the material to be 97% pure; ret. time = 12.6 min.

Example (Compound 122)



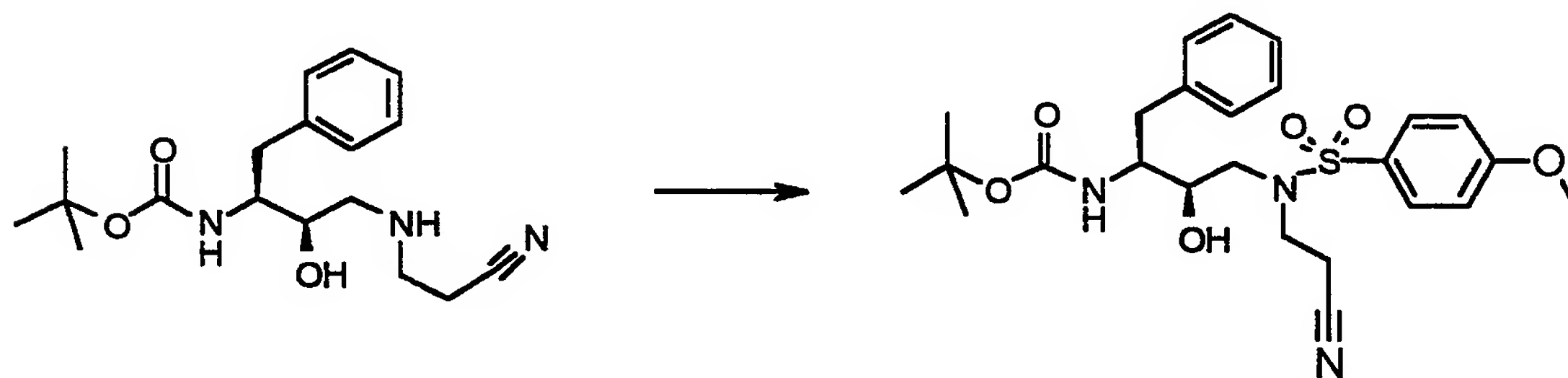
- 20 [(2*R*,3*S*)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino-2-cyanoethyl) butane].

[(2*R*,3*S*)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1- amino) butane] (0.503 g, 1.8 mmol) was

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dissolved in EtOH (6 mL) and cooled to 0 °C. This was followed by the addition of acrylonitrile (118 uL, 1.8 mmol). The reaction was stirred under N₂ for 15 hrs and gradually warmed to room temperature as the ice bath
5 melted. The reaction mixture was then concentrated in vacuo and purified by silica gel chromatography (gradient: 2 % MeOH/CH₂Cl₂ to 4% MeOH/CH₂Cl₂). Isolated 339 mg of the mono-addition product (56% yield).

Example (Compound 123)



10

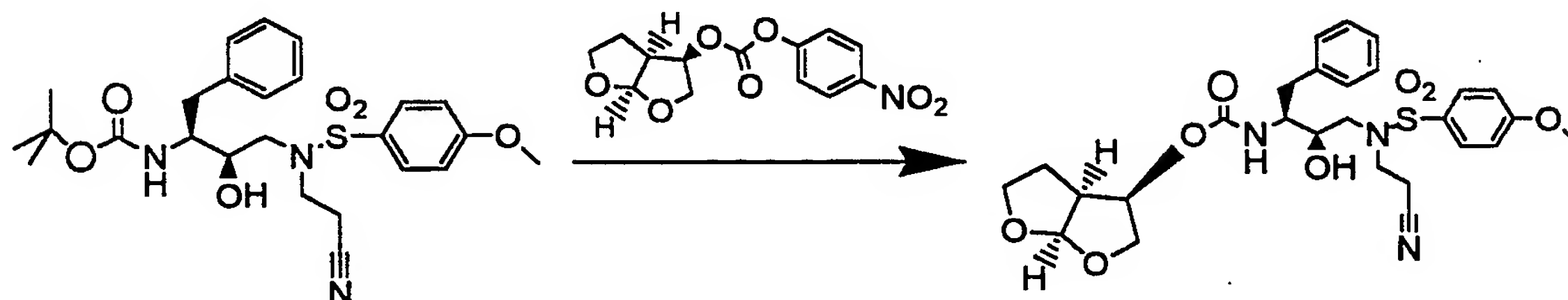
[(2R,3S) -3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-
(1-amino-(2-cyanoethyl (4-methoxyphenylsulfonyl))
butane].

[(2R,3S) -3-*tert*-Butoxycarbonylamino-2-hydroxy-
15 4-phenyl-(1-amino-2-cyanoethyl) butane]. (124 mg, 0.4
mmol) was combined with 4-methoxyphenylsulfonyl chloride
(146 mg, 0.7 mmol) in DMF and cooled to 0 °C. This was
followed by the addition of diisopropyl ethyl amine (186
uL, 1.1 mmol). The reaction was then warmed to room
20 temperature and stirred for 15 hours. The reaction was
quenched by the addition of 3 mL of saturated NaHCO₃,
followed by washing with sat. NaHCO₃, KHSO₃ and then
brine. The organic phase was dried with MgSO₄ and the
solvent was removed *in vacuo*. Purification by SiO₂ (1:3,
25 EtOAc:Hex to 2:1 EtOAc:Hex). Isolated 0.038 g of the

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product as a colorless oil: 20.8 % yield, $R_f=0.27$ (1:1 EtOAc: Hex/SiO₂), MS(ES) = 521 amu (M+1).

Example (Compound 124)



{ (1S,2R)-3-[(4-methoxybenzenesulfonyl)-(2-cyanoethyl)-
5 amino]-1-benzyl-2-hydroxy-propyl}-carbamic acid
(3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

Step 1:

[(2R,3S)-3-tert-Butoxycarbonylamino-2-hydroxy-
4-phenyl-(1-amino-(2-cyanoethyl (4-
10 methoxyphenylsulfonyl)) butane] (521 mg, 0.1 mmol) was
dissolved in 1 mL of CH₂Cl₂ followed by the addition of 1
mL of TFA. The reaction was stirred at room temperature
for 15 minutes followed by concentration *in vacuo* to
provide 655 mg of the desired deprotected amine. This
15 material was used without further purification.

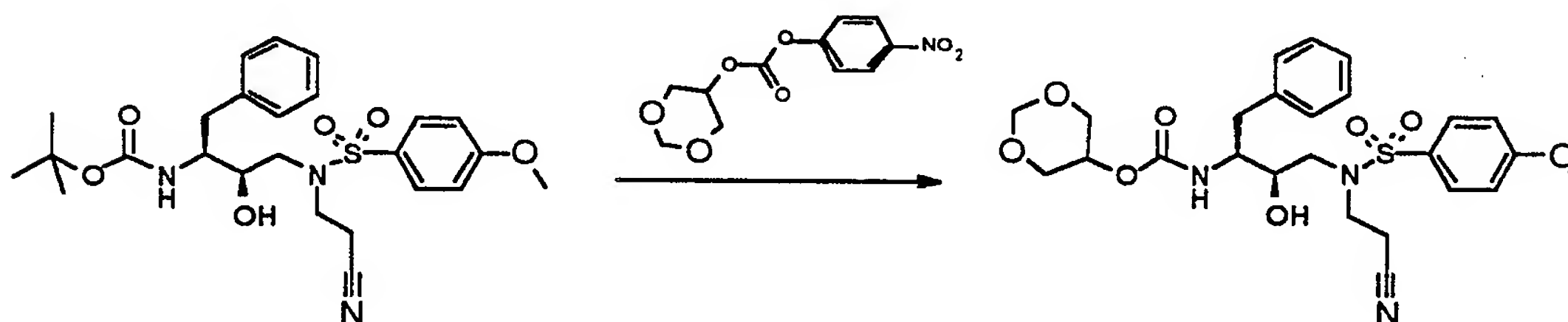
Step 2:

31.2 mg (0.1 mmol) of the material provided in
the above deprotection step was then dissolved into DMF
(1 mL) followed by the addition of the
20 3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl mixed
carbonate (30.5 mg, 0.1 mmol). After cooling to 0 °C
diisopropylethyl amine was added and the reaction was
stirred for 3 hours while gradually warming to room
temperature. The reaction mixture was then diluted with

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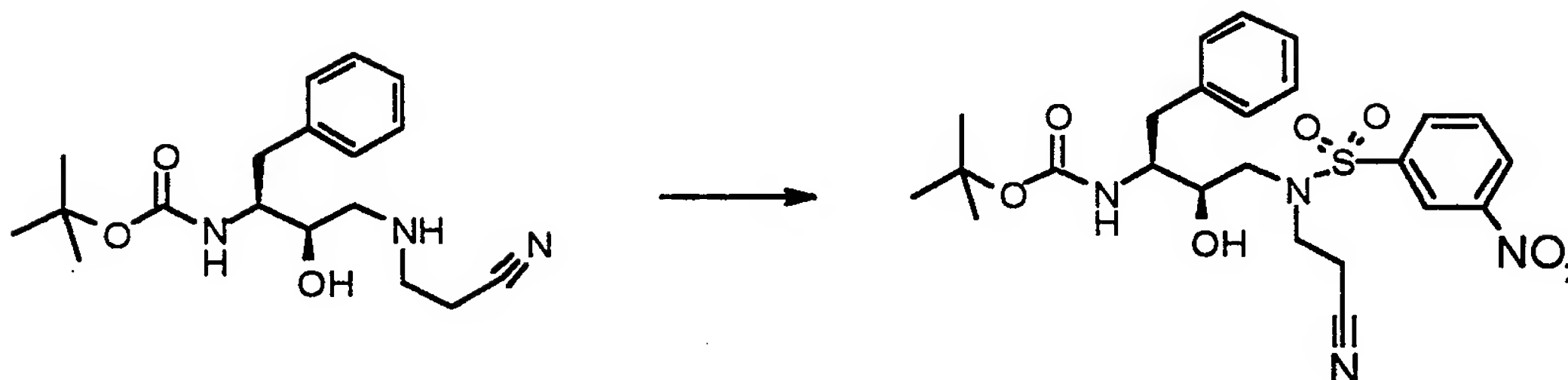
EtOAc washed with sat. NaHCO_3 , KHSO_3 and then brine. The organic phase was dried with MgSO_4 and the solvent was removed *in vacuo*. Purification by Prep. TLC (SiO_2 2:1, EtOAc:Hex). Isolated 0.0217 g of the product as a white solid: $R_f=0.28$ (2:1 EtOAc: Hex, / SiO_2), MS(ES) = 560 amu (M+1).

Example (Compound 125)



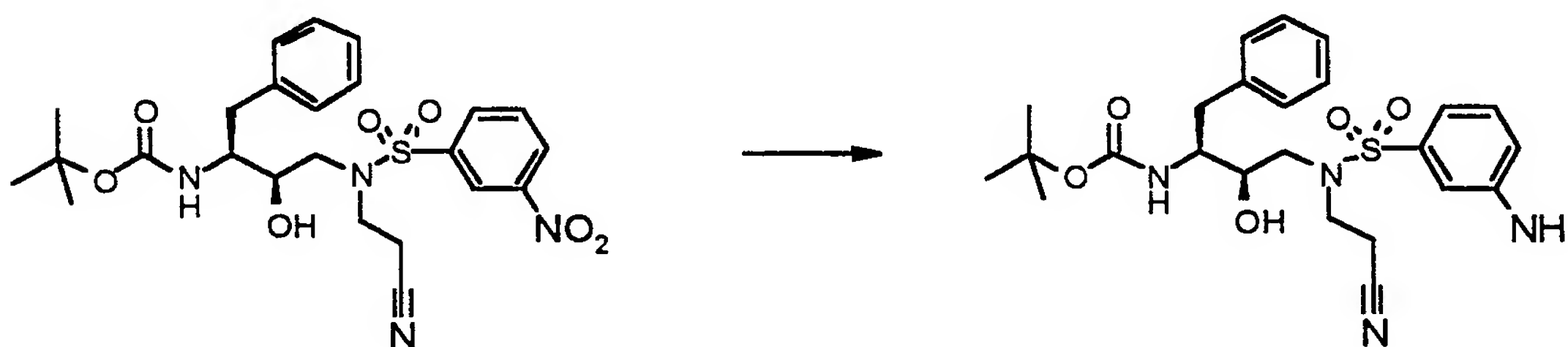
34.3 mg (0.1 mmol) of the material provided in the above deprotection step (Step 1, Example (Compound 122)) was then dissolved into DMF (1 mL) followed by the 1,3-dioxan-5-yl derived mixed carbonate (127.9 mg, 0.1 mmol). After cooling to 0 °C, diisopropylethylamine was added and the reaction was stirred for 15 hours (warmed to room temperature overnight). The reaction mixture was then diluted with EtOAc washed with sat. NaHCO_3 , KHSO_3 and then brine. The organic phase was dried with MgSO_4 and the solvent was removed *in vacuo*. Purification by Prep. TLC (SiO_2 2:1, EtOAc:Hex). Isolated 0.0122 g of the product as a white solid: $R_f=0.34$ (2:1, EtOAc:Hex/ SiO_2), MS(ES) = 534 amu (M+1).

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Example (Compound 126)

[(2R,3S)-3-tert-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino-(2-cyanoethyl (3-nitrophenylsulfonyl)) butane].

5 [(2R,3S)-3-tert-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino-2-cyanoethyl) butane]. (100.6 mg, 0.3 mmol) was combined with 3-nitrophenylsulfonyl chloride (134 mg, 0.6 mmol) in DMF and cooled to 0 °C. This was followed by the addition of diisopropyl ethyl amine (160
10 uL, 0.9 mmol). The reaction was then warmed to room temperature and stirred for 15 hours. The reaction was quenched by the addition of 3 mLs of saturated NaHCO₃, followed by washing with sat. NaHCO₃, KHSO₃ and then brine. The organic phase was dried with MgSO₄ filtered
15 and the solvent was removed *in vacuo*. Purification by SiO₂ (1:3, EtOAc:Hex to 2:1 EtOAc:Hex). Isolated 0.0389 g of the product as a colorless oil: 24.9 % yield, R_f=0.39 (1:1 EtOAc: Hex/SiO₂).

Example (Compound 127)

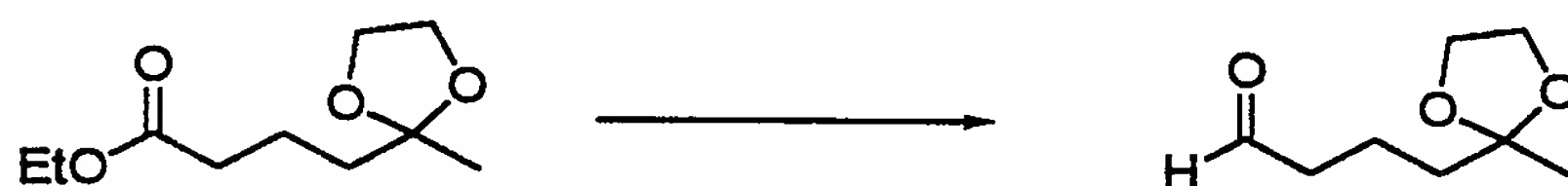
-152-

[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino-(2-cyanoethyl (3-aminophenylsulfonyl)) butane].

[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino-(2-cyanoethyl (3-nitrophenylsulfonyl)) butane] (4.0 mg) was dissolved in methanol (1 mL) followed by catalytic 10% Pd/C. The reaction vessel was purged of air and pressurized with H₂ at ~1 ATM and stirred for 3 hours. The reaction was then filtered and concentrated *in vacuo* to provide 2.1 mg of a colorless film: R_f=0.22 (1:1 EtOAc: Hex, /SiO₂), MS(ES) = 489 amu (M+1).

Example (Compound 201)

Step 1:



15 4-(2-Methyl-1,3-dioxolan-2-yl)butanal

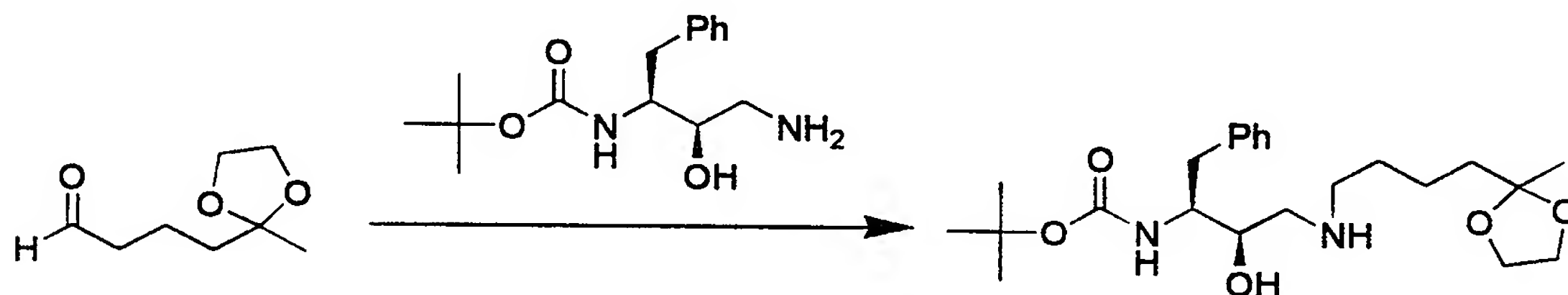
A stirred slurry of lithium aluminum hydride (940 mg, 24.7 mmol) in anhydrous ether (25 mL) was cooled in an ice bath. Ethyl 4-(2-methyl-1,3-dioxolan-2-yl)butanoate [Benchikh-le-Hocine, M.; Do Khac, D.; Fetizon, M.; Prange, T. *Synthetic Communications* 1992, 22, 1871-1882] (5.0 g, 24.7 mmol) in ether (25 mL) was added dropwise over 20 minutes. The reaction was stirred at 5 °C for 2 hours and then quenched by the slow addition of water (1 mL), followed by 15% sodium hydroxide (1 mL) and water (3 mL). After stirring for 30 minutes, the resulting precipitate was filtered and rinsed with ether. The ether was dried (magnesium sulfate) and concentrated *in vacuo* to afford the desired alcohol (3.6 g, 90%) as a colorless liquid.

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¹H NMR (CDCl₃): δ 1.30 (3H, s), 1.42-1.72 (7H, m), 3.64 (2H, t), 3.88-3.98 (4H, m).

To a stirred suspension of pyridinium chlorochromate (7.1 g, 32.8 mmol) in dichloromethane (35 mL) was added a solution of the above alcohol (3.5 g, 21.8 mmol) in dichloromethane (15 mL). After stirring at ambient temperature for 2 hours, the reaction was diluted with ether (50 mL) and filtered through celite. The filtrate was concentrated in vacuo and chromatographed (silica gel, ethyl acetate/hexane, 1:4) to afford the title compound (1.8 g, 50%) as a clear liquid; ¹H NMR (CDCl₃): δ 1.30 (3H, s), 1.62-1.78 (4H, m), 2.46 (2H, t), 3.88-3.98 (4H, m), 9.78 (1H, s).

Step 2:



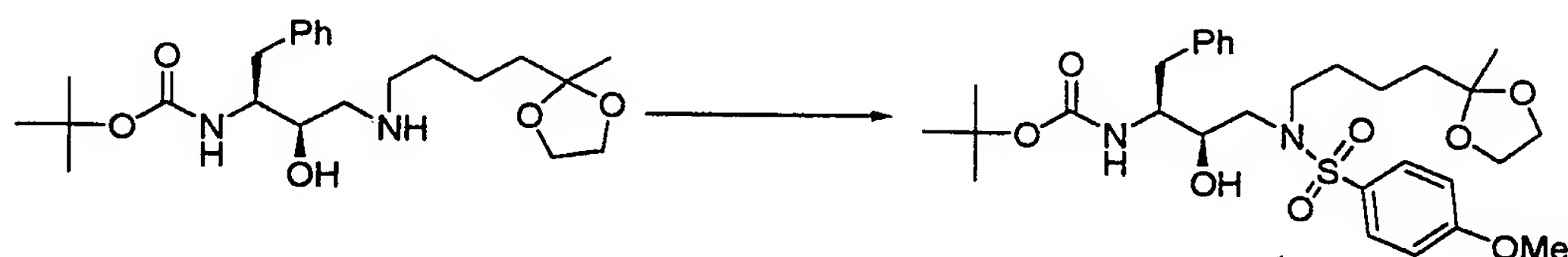
tert-Butyl N-((1S,2R)-1-benzyl-2-hydroxy-3-[4-(2-methyl-1,3-dioxolan-2-yl)butyl] aminopropyl)carbamate

To a suspension of the product of Step 1 (1.8 g, 11.4 mmol) and tert-butyl N-(1S,2R)-3-amino-1-benzyl-2-hydroxypropylcarbamate (3.2 g, 11.4 mmol) in 1,2-dichloroethane (30 mL), tetrahydrofuran (30 mL) and N,N-dimethylformamide (20 mL) was added sodium triacetoxy borohydride (3.6 g, 17.0 mmol) and the mixture was stirred under nitrogen atmosphere for 3.5 hours. The mixture was filtered and the filtrate was concentrated to 25 mL. The residue was partitioned between ethyl acetate and 2M sodium hydroxide/water. The organic phase was

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washed with water, brine, dried (sodium sulfate), and evaporated. The residue was chromatographed (silica gel, dichloromethane/methanol, 95:5) to provide the title compound as a white solid.

5 Step 3:



tert-Butyl *N*-((1*S*,2*R*)-1-benzyl-2-hydroxy-3-[(4-methoxyphenyl)sulfonyl][4-(2-methyl-1,3-dioxolan-2-yl)butyl]aminopropyl)carbamate

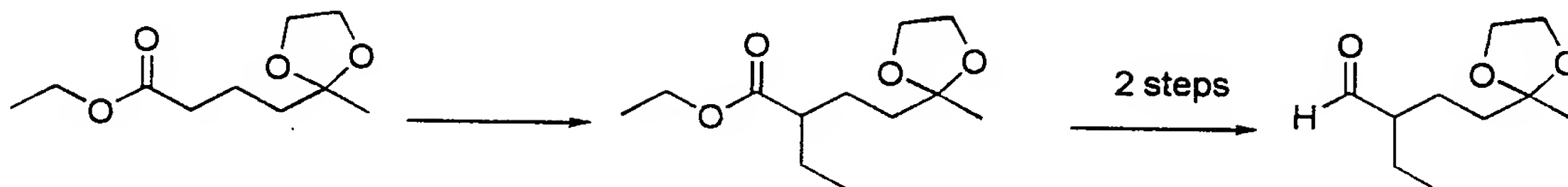
10 To a solution of the product of Step 2 (0.63 g, 1.49 mmol) in anhydrous dichloromethane (7 mL) at 0°C was added *N*-ethyldiisopropylamine (0.38 g, 0.51 mL, 2.98 mmol) and a solution of 4-methoxybenzenesulfonyl chloride (0.32 g, 1.56 mmol) in anhydrous dichloromethane (2 mL)
 15 and the mixture was stirred at ambient temperature under nitrogen atmosphere for 18 hours. The mixture was washed sequentially with 1M hydrochloric acid, sodium bicarbonate/water, brine, dried (sodium sulfate) and concentrated. The residue was chromatographed (silica
 20 gel, dichloromethane/methanol, 97:3) to provide the title compound as a thick oil.

¹H NMR (DMSO-*d*₆): δ 1.18 (3H, s), 1.21 (9H, s), 1.4-1.5 (4H, m), 2.45 (2H, brs), 2.82 (1H, dd), 2.94-3.00 (2H, m), 3.18 (1H, dt), 3.28-3.34 (2H, m), 3.45-3.60 (2H, m),
 25 3.75-3.84 (7H, m), 5.0 (1H, brs), 6.7 (1H, d), 7.08 (2H, d), 7.10-7.25 (5H, m), 7.7 (2H, d); MS: 615.1 (MH⁺)

Example (Compound 202)

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Step 1:

**2-Ethyl-4-(2-methyl-1,3-dioxolan-2-yl)butanal**

To a freshly prepared solution of lithium diisopropylamide (40.0 mmol in tetrahydrofuran/hexane; 50 mL, 1:1) at -78°C was added a solution of ethyl-4-(2-methyl-1,3-dioxolan-2-yl) butanoate (6.7 g, 33.3 mmol) in tetrahydrofuran (15 mL) over 15 minutes. After stirring for 15 minutes, ethyl iodide (4.0 mL, 50.0 mmol) was added rapidly via syringe. The reaction was stirred at -78°C for 6 hours and then allowed to warm slowly to ambient temperature and stir for 16 hours. The reaction was quenched by the addition of water (50 mL) and extracted with ether (3 x 50 mL). The combined organics were washed with brine (50 mL), dried (magnesium sulfate), concentrated *in vacuo* and chromatographed (silica gel, ethyl acetate/hexane, 1:4) to afford ethyl 2-ethyl-4-(2-methyl-1,3-dioxolan-2-yl) butanoate (2.6 g, 33%) as a colorless liquid.

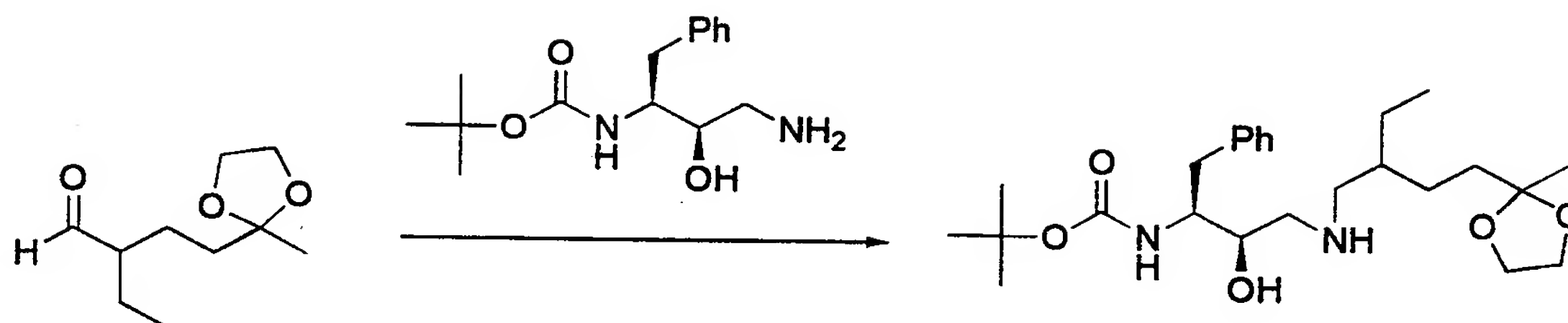
^1H NMR (CDCl_3) : δ 0.88 (3H, t), 1.23 (3H, t), 1.30 (3H, s), 1.45-1.75 (6H, m), 2.20-2.28 (1H, m), 3.85-3.95 (4H, m), 4.12 (2H, q).

The above ester (2.5g, 10.9mmol) was subjected to the conditions used in Step 1 (Example (Compound 201)) to afford the title compound (1.2 g, 60%) as a clear liquid.

^1H NMR (CDCl_3): δ 0.90 (3H, t), 1.30 (3H, s), 1.45-1.78 (6H, m), 2.14-2.22 (1H, m), 3.87-3.97 (4H, m), 9.58 (1H, s).

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Step 2:



tert-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-[2-ethyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl]amino-2-hydroxypropyl) carbamate

The product from Step 1 was subjected to
 5 procedure Step 2, Example (Compound 201) to afford the
 title compound as a white solid (mixture of diastereomers
 by ¹H NMR spectroscopy);

¹H NMR (DMSO-*d*₆): δ 0.78 (3H, t), 1.17 (3H, s), 1.20-1.35
 (5H, m), 1.22 (9H, s), 1.48 (2H, t), 2.33-2.58 (5H, m),
 10 2.92-2.98 (1H, m), 3.34-3.55 (2H, m), 3.75-3.84 (4H, m),
 4.78 (1H, br s), 6.68 (1H, dd), 7.05-7.25 (5H, m) ; MS:
 451 (MH⁺).

Step 3:



tert-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-[2-ethyl-4-(2-methyl-1,3-
 15 dioxolan-2-yl)butyl][(4-methoxyphenyl)sulfonyl]amino-2-
 hydroxypropyl) carbamate

The product from above Step 2 was subjected to
 the procedure outlined in Step 3 of Example (Compound

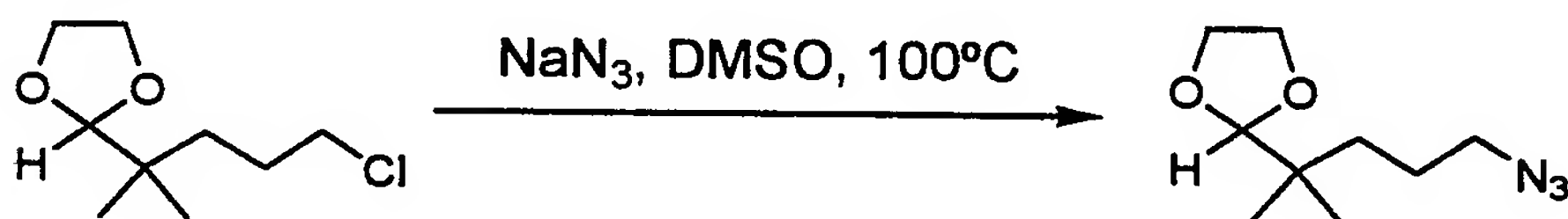
-157-

201) to give the title compound as a pale yellow glass (mixture of diastereomers by ^1H NMR spectroscopy);

^1H NMR ($\text{DMSO}-d_6$): δ 0.79 (3H, td), 1.2 (3H, s), 1.24 (9H, s), 1.3-1.6 (3H, m), 1.7 (1H, brs), 2.45-2.55 (2H, m),
5 2.73 (1H, dd), 2.85 (1H, dd), 2.9 (1H, d), 3.1 (1H, td), 3.40-3.65 (4H, m), 3.75-3.90 (7H, m), 4.96 (1H, brs), 6.7 (1H, d), 7.05 (2H, d), 7.1-7.3 (5H, m), 7.78 (2H, d); MS: 643.1 (M + 23)

Example (Compound 203)

10 Step 1:



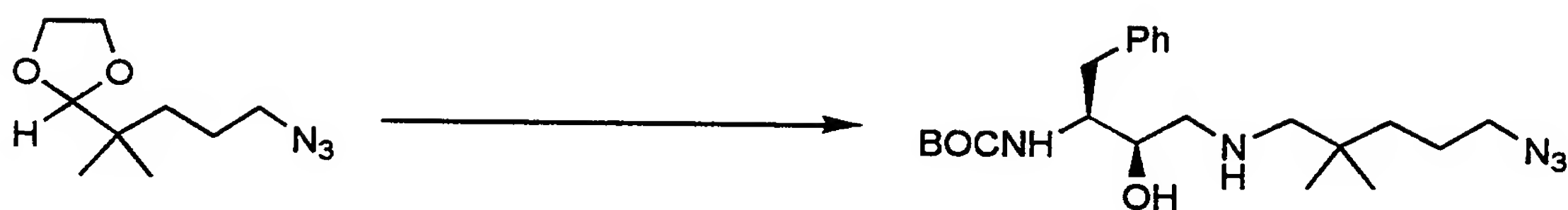
4-(1,3-dioxolan-2-yl)-4-methylpentyl azide

A solution of 0.74 g (3.8 mmol) of 2-(4-chloro-1,1-dimethylbutyl)-1,3-dioxolane and 0.50 g (7.7 mmol) of
15 sodium azide in 5 mL of DMSO was heated to 100°C with stirring. After 18 hours the solution was cooled to RT, diluted with water, and the mixture extracted with ether (3x). The combined ether extracts were washed with water (3x), dried over MgSO_4 , and concentrated in vacuo to
20 afford 0.69 g (90%) of the desired compound as a light yellow liquid.

^1H NMR (CDCl_3): δ 4.60 (1H), 3.91 (4H), 3.29 (2H), 1.67 (2H), 1.41 (2H), 0.96 (6H).

Step 2:

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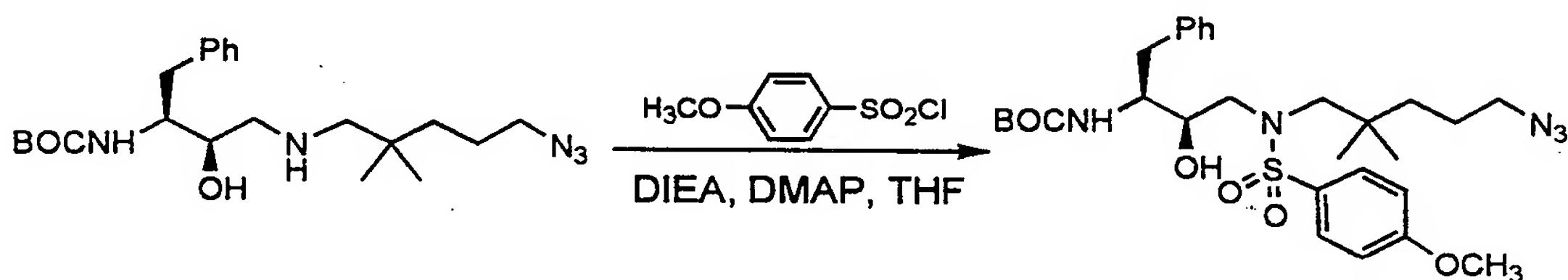
tert-butyl *N*-(1*S*,2*R*)-3-[(5-azido-2,2-dimethylpentyl)amino]-1-benzyl-2-hydroxypropylcarbamate

A solution of 0.44 g (2.2 mmol) of 4-(1,3-dioxolan-2-yl)-4-methylpentyl azide in 15 mL of THF was
5 treated with 3 mL of 1M aqueous HCl and the solution stirred at reflux. After 4 hours the solution was cooled to RT and poured into 40 mL of rapidly stirred saturated aqueous NaHCO₃. The resulting mixture was extracted with ether (3x). The combined ether extracts were washed with
10 water (2x), dried over MgSO₄ and concentrated to a volume of approximately 3 mL. This solution was added to a solution of 0.62 g (2.2 mmol) of *tert*-butyl *N*-(1*S*,2*R*)-3-amino-1-benzyl-2-hydroxypropylcarbamate in 20 mL of 1:1 THF/DMF. The solution was treated with 1 g of powdered
15 4A molecular sieves and the mixture heated at 50°C. After 2 hours the mixture was cooled in an ice water bath and treated with 0.47 g (2.2 mmol) of NaBH(OAc)₃. After stirring at RT for 18 hours the mixture was concentrated *in vacuo*. The residue was suspended in CH₂Cl₂ and stirred
20 vigorously with 0.5M aqueous NaOH for 5 minutes. The layers were separated and the aqueous phase extracted with two additional portions of CH₂Cl₂. The combined CH₂Cl₂ solutions were washed with water (2x), dried over MgSO₄, and concentrated to give a clear viscous oil. This
25 material was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.45 g (49%) of the desired product as a white foam.

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$^1\text{H-NMR}$ (DMSO-d_6): δ 7.27-7.05 (5H), 6.70 (1H), 4.73 (1H), 3.49 (1H), 3.39 (1H), 3.25 (2H), 2.92 (1H), 2.60-2.42 (4H), 2.22 (2H), 1.41 (3H), 1.21 (10H), 0.80 (6H).
 MS (ESI): 420 (M+H).

5 Step 3:



tert-butyl *N*-((1*S*,2*R*)-3-(5-azido-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl) carbamate

A solution of 0.40 g (0.95 mmol) of *tert*-butyl *N*-(1*S*,2*R*)-3-[(5-azido-2,2-dimethylpentyl)amino]-1-benzyl-2-hydroxypropylcarbamate in 10 mL of anhydrous THF was treated with 0.20 g (0.95 mmol) of 4-methoxybenzenesulfonyl chloride, 0.17 mL (0.95 mmol) of *N,N*-diisopropylethylamine, and 35 mg (0.29 mmol) of DMAP. The resulting solution was stirred at RT. After 2 hours thin layer chromatography ("tlc") (SiO_2 , hexane/EtOAc) indicated reaction to be complete. The solution was concentrated in the presence of silica gel and the material subjected to flash chromatography (SiO_2 , 8:2 to 7:3 hexane/EtOAc) to afford 0.50 g (89%) of the desired compound as a viscous oil.

$^1\text{H-NMR}$ (DMSO-d_6): δ 7.76 (2H), 7.30-7.03 (7H), 6.65 (1H), 5.01 (1H), 3.85 (3H), 3.74 (1H), 3.48-3.23 (4H), 2.95 (2H), 2.81 (1H), 2.50 (2H), 1.52 (2H), 1.40-1.20 (11H), 0.94 (6H); MS (ESI): 590 (M+H).

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Example (Compound 204)

Step 1:

**4-(2-Methyl-1,3-dioxolan-2-yl)butanal**

A stirred slurry of lithium aluminum hydride (940 mg, 24.7 mmol) in anhydrous ether (25 mL) was cooled in an ice bath. Ethyl 4-(2-methyl-1,3-dioxolan-2-yl)butanoate [Benchikh-le-Hocine, M.; Do Khac, D.; Fetizon, M.; Prange, T. *Synthetic Communications* **1992**, 22, 1871-1882] (5.0 g, 24.7 mmol) in ether (25 mL) was added dropwise over 20 minutes. The reaction was stirred at 5 °C for 2 hours and then quenched by the slow addition of water (1 mL), followed by 15% sodium hydroxide (1 mL) and water (3 mL). After stirring for 30 minutes, the resulting precipitate was filtered and rinsed with ether. The ether was dried (magnesium sulfate) and concentrated *in vacuo* to afford the desired alcohol (3.6 g, 90%) as a colorless liquid;

¹H NMR (CDCl₃): δ 1.30 (3H, s), 1.42-1.72 (7H, m), 3.64 (2H, t), 3.88-3.98 (4H, m);

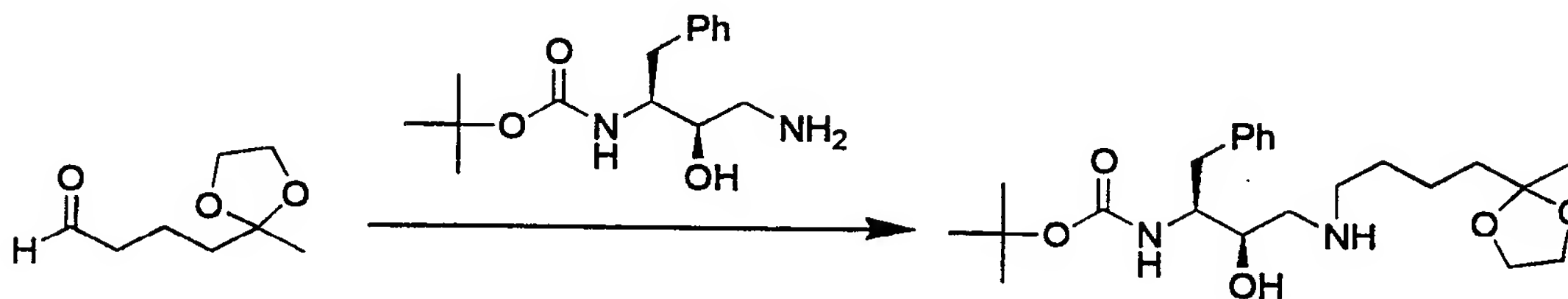
To a stirred suspension of pyridinium chlorochromate (7.1 g, 32.8 mmol) in dichloromethane (35 mL) was added a solution of the above alcohol (3.5 g, 21.8 mmol) in dichloromethane (15 mL). After stirring at ambient temperature for 2 hours, the reaction was diluted with ether (50 mL) and filtered through celite. The filtrate was concentrated *in vacuo* and chromatographed

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(silica gel, ethyl acetate/hexane, 1:4) to afford the title compound (1.8 g, 50%) as a clear liquid;

^1H NMR (CDCl_3): δ 1.30 (3H, s), 1.62-1.78 (4H, m), 2.46 (2H, t), 3.88-3.98 (4H, m), 9.78 (1H, s).

5 Step 2:

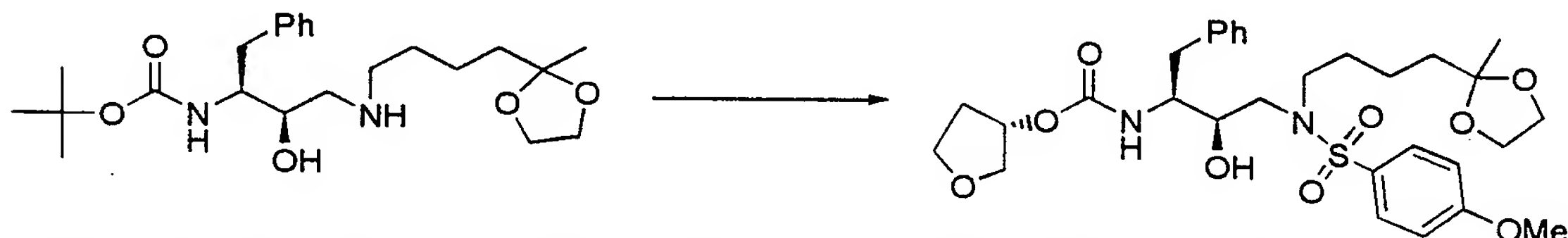


tert-Butyl *N*-((1*S*,2*R*)-1-benzyl-2-hydroxy-3-[4-(2-methyl-1,3-dioxolan-2-yl)butyl] aminopropyl)carbamate

The product from step a was subjected to the procedure used in Step 2 of Example (Compound 201) to afford the title compound as a white solid;

^1H NMR ($\text{DMSO}-d_6$): δ 1.18 (3H, s), 1.23 (9H, s), 1.26-1.41 (4H, m), 1.48-1.55 (2H, m), 1.98 (1H, brs), 2.39-2.59 (5H, m), 2.95 (1H, dd), 3.34-3.54 (2H, m), 3.75-3.84 (4H, m), 4.78 (1H, brs), 6.65 (1H, d), 7.08-7.27 (5H, m); MS: 423 (MH^+).

Step 3:



tert-Butyl *N*-((1*S*,2*R*)-1-benzyl-2-hydroxy-3-[4-(4-methoxyphenyl)sulfonyl][4-(2-methyl-1,3-dioxolan-2-yl)butyl]aminopropyl)carbamate

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The product from Step 2 was subjected to the procedure used in step 3 of Example (Compound 201) to give the title compound as a thick oil;

¹H NMR (DMSO-d₆): δ 1.18 (3H, s), 1.21 (9H, s), 1.4-1.5 (4H, m), 2.45 (2H, brs), 2.82 (1H, dd), 2.94-3.00 (2H, m), 3.18 (1H, dt), 3.28-3.34 (2H, m), 3.45-3.60 (2H, m), 3.75-3.84 (7H, m), 5.0 (1H, brs), 6.7 (1H, d), 7.08 (2H, d), 7.10-7.25 (5H, m), 7.7 (2H, d); MS: 615.1 (MH⁺).

Example (Compound 205)



10 **(3S)-Tetrahydro-3-furanyl N-(1S,2R)-1-benzyl-2-hydroxy-3-[[(4-methoxyphenyl) sulfonyl] (5-oxohexyl) amino]-propylcarbamate**

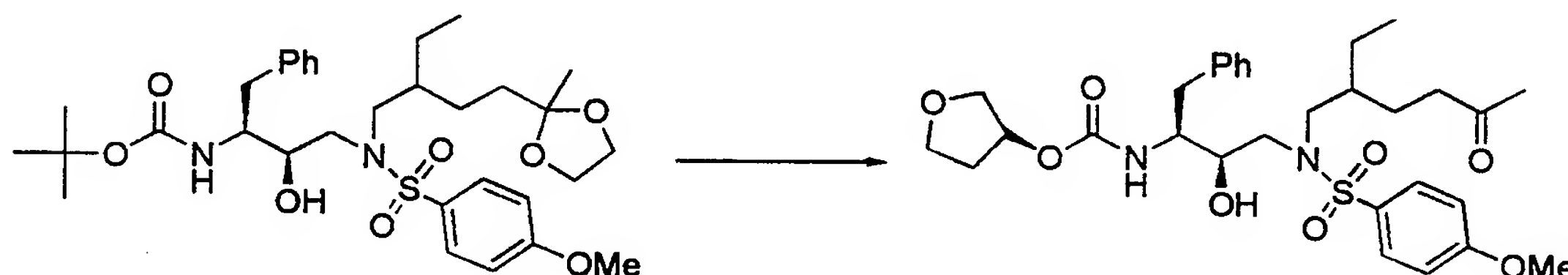
The product from Step 3 of Example (Compound 204) (0.22 g, 0.4 mmol) was dissolved in dichloromethane (2.5 mL) at 0°C was added trifluoroacetic acid (2.5 mL) and a drop of water, and the mixture was stirred at ambient temperature for 1 hour. Solvent was evaporated and coevaporated with dichloromethane. The residue was dissolved in dichloromethane and washed with 0.1 M sodium hydroxide/water, water, brine, dried (sodium sulfate), concentrated and dried in vacuo. The resulting pale yellow foam was dissolved in acetonitrile (4 mL) and N-ethyldiisopropylamine (0.08 mL, 0.059 g, 0.46 mmol) and N-succinimidyl (3S)-tetrahydro-3-furanyl carbonate (0.098 g, 0.43 mmol) were added. The mixture was stirred at ambient temperature for 1 hour and the solvent was

-163-

evaporated. The residue was chromatographed (silica gel, dichloromethane/methanol, 98:2) to give the title compound as a white foam;

¹H NMR (DMSO-d₆): δ 1.3-1.5 (4H, m), 1.73 (1H, brs), 2.05 (4H, brs), 2.35 (2H, brs), 2.5 (1H, brs), 2.8 (1H, dd), 2.95 (2H, brd), 3.15-3.20 (1H, m), 3.35 (H, t), 3.45-3.75 (5H, m), 3.8 (3H, s), 4.93 (1H, s), 5.1 (1H, brs), 7.1-7.3 (8H, m), 7.7 (2H, d); MS: 563.1 (MH⁺).

Example (Compound 206)



10 **(3S)-Tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-(2-ethyl-5-oxohexyl) [(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate**

The product from Step 3 of Example (Compound 202) was subjected to the procedure used in Example
15 (Compound 205) to provide the title compound as a thick oil (mixture of diastereomers by ¹H NMR spectroscopy);

¹H NMR (DMSO-d₆): δ 0.85 (3H, td), 1.2-1.3 (2H, m), 1.4-1.6 (3H, m), 1.7-1.9 (2H, m), 2.10-2.15 (1H, m), 2.18 (3H, s), 2.2-2.3 (2H, m), 2.85 (1H, d), 2.90 (1H, d),
20 3.05 (1H, d), 3.15 (1H, dd), 3.38-3.42 (1H, m), 3.45-3.80 (6H, m), 3.9 (3H, s), 5.0 (1H, brt), 5.1 (1H, t), 7.1 (2H, d), 7.2-7.4 (6H, m), 7.8 (2H, m): MS: 591.1 (MH⁺).

Example (Compound 207)

Step 1:

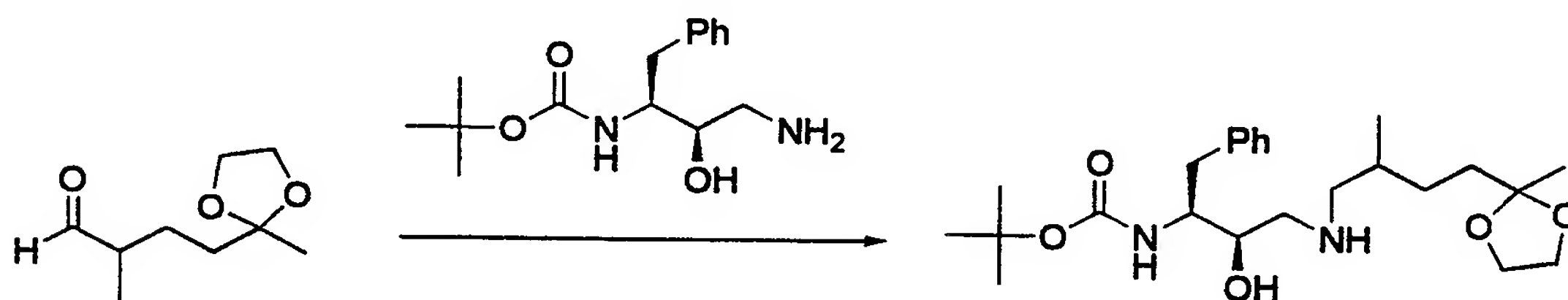
-164-

**2-Methyl-4-(2-methyl-1,3-dioxolan-2-yl)butanal**

The title compound was prepared by the procedure used in Example (Compound 202) using methyl iodide in the alkylation step.

^1H NMR (CDCl_3): δ 1.10 (3H, d), 1.31 (3H, s), 1.40-1.88 (4H, m), 2.26-2.40 (1H, m), 3.86-3.98 (4H, m), 9.61 (1H,

Step 2:



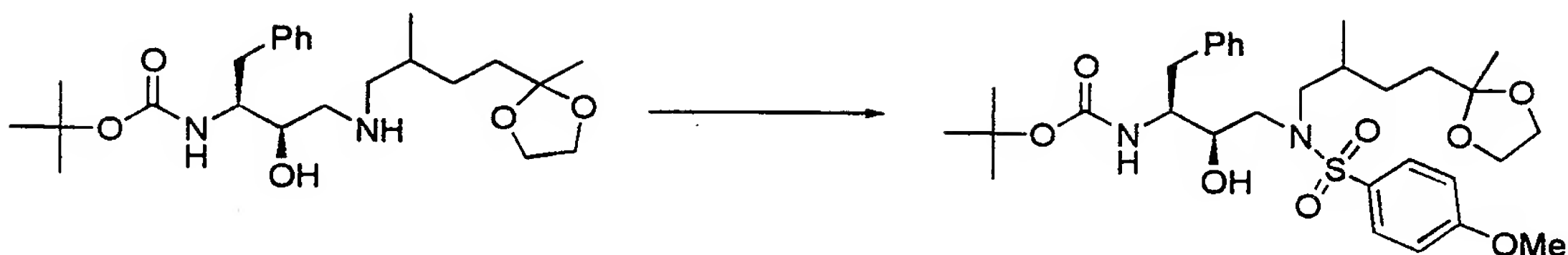
tert-Butyl N-((1S,2R)-1-benzyl-2-hydroxy-3-[2-methyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl]aminopropyl)carbamate

The product from Step 1 was subjected to procedure used in Step 2 of Example (Compound 201) to provide the title compound as a white solid (mixture of diastereomers by ^1H NMR spectroscopy);

^1H NMR ($\text{DMSO}-d_6$): δ 0.82 (3H, d), 1.18 (3H, s), 1.23 (9H, s), 1.33-1.62 (5H, m), 1.84 (1H, br), 2.23-2.58 (5H, m), 2.96 (1H, d), 3.36-3.55 (2H, m), 3.75-3.85 (4H, m), 4.78 (1H, br s), 6.67 (1H, dd), 7.07-7.25 (5H, m); MS: 437 (MH $^+$).

-165-

Step 3:



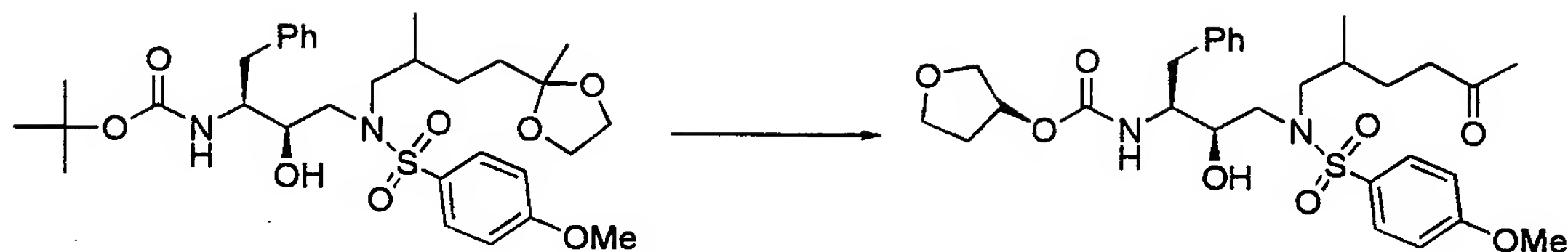
tert-Butyl *N*-((1*S*,2*R*)-1-benzyl-2-hydroxy-3-[2-methyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl]amino)propyl carbamate
 5 (4-methoxyphenyl)sulfonyl]aminopropyl) carbamate

The product from Step 2 was subjected to Step 3 of Example (Compound 201) to afford the title compound as a yellow foam (mixture of diastereomers by ^1H NMR spectroscopy);

10 ^1H NMR ($\text{DMSO}-d_6$): δ 0.8 (3H, d), 1.00-1.05 (1H, m), 1.1 (1H, brs), 1.2 (9H, s), 1.2-1.3 (1H, m), 1.35-1.50 (2H, m), 1.6 (1H, td), 1.75-1.83 (1H, m), 2.5 (1H, dd), 2.8 (1H, dd), 2.85 (1H, dd), 2.93 (1H, dd), 3.05 (1H, dd), 3.15-3.2 (2H, m), 3.37-3.46 (1H, m), 3.59 (1H, brs),
 15 3.80-3.95 (7H, m), 4.95 (1H, brs), 6.7 (1H, d), 7.1 (2H, d), 7.15-7.24 (5H, m), 7.75 (2H, d); MS: 629.1 ($M + 23$).

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Step 4:



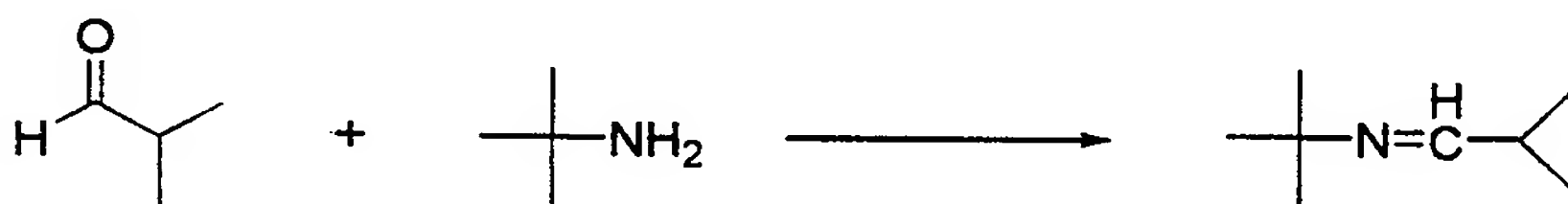
(3S)-Tetrahydro-3-furanyl N-(1S,2R)-1-benzyl-2-hydroxy-3-[[(4-methoxyphenyl) sulfonyl] (2-methyl-5-oxohexyl) amino] propylcarbamate

5 The product from Step 3 was subjected to the procedure in Example (Compound 205) to give the title compound as a white foam (mixture of diastereomers by ^1H NMR spectroscopy);

^1H NMR (DMSO- d_6): δ 0.83 (3H, d), 1.2-1.3 (2H, m), 1.5-1.6 (1H, m), 1.8-1.9 (2H, m), 2.05-2.10 (1H, m), 2.2 (3H, s), 2.3-2.5 (2H, m), 2.8 (1H, dd), 2.9 (1H, brs), 3.05 (1H, brd), 3.15 (1H, dd), 3.30-3.45 (2H, m), 3.5-3.8 (6H, m), 3.9 (3H, s), 5.0 (1H, brt), 5.1 (1H, brdd), 6.2 (2H, d), 6.25-6.40 (6H, m), 7.8 (2H, m); MS: 577.5 (MH^+).

Example (Compound 208)

Step 1:



N-(2-methylpropylidene)-2-methyl-2-propanamine

To a stirred solution of *t*-butylamine (20.0 mL, 190 mmol) in anhydrous ether (200 mL) was added magnesium sulfate (20 g). A solution of isobutyraldehyde (17.3 mL,

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190 mmol) in ether (50 mL) was then added dropwise over 30 minutes. After stirring at ambient temperature for 3 days, the reaction was filtered and the ether removed in vacuo (300 mm). The residual liquid was distilled to afford the title compound (14.0g, 58%) as a colorless liquid; bp 52 °C (75 mm);

¹H NMR (CDCl₃): δ 1.01 (6H, d), 1.13 (9H, s), 2.37 (1H, m), 7.36 (1H, d).

2,2-Dimethyl-4-(2-methyl-1,3-dioxolan-2-yl)butanal

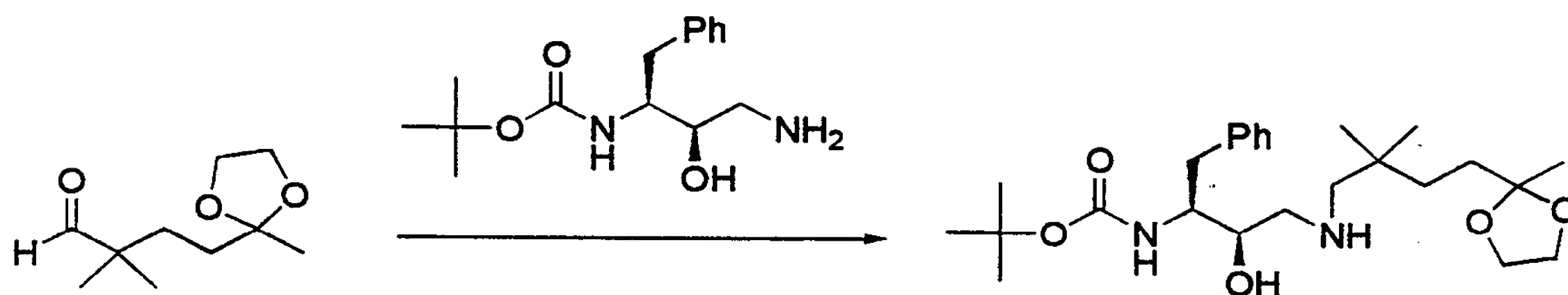
10 A freshly prepared solution of lithium diisopropylamide, was generated by the addition of *n*-butyllithium (16.2 mL, 25.9 mmol, 1.6 M in hexane) to a solution of diisopropylamine (6.6 mL, 47.0 mmol) in tetrahydrofuran (20 mL) at -20°C, and allowed to warm to 0
15 °C and stir for 30 minutes. *N*-(2-methylpropylidene)-2-methyl-2-propanamine (3.0 g, 23.5 mmol) in tetrahydrofuran (10 mL) was added dropwise over 10 minutes. After stirring at 0°C for 2 hours, 2-bromoethyl-2-methyl-1,3-dioxolane (5.0 g, 25.9 mmol) in
20 tetrahydrofuran (10 mL) was added over 15 minutes. The reaction was allowed to warm to ambient temperature and stirring was continued for 16 hours. After cooling in an ice bath, an aqueous solution of oxalic acid (100 mL, 10 wgt %) was added and reaction stirred for 1 hour. The
25 layers were separated and the aqueous was extracted with ether (100 mL). The combined organic layers were washed with brine (50 mL), saturated sodium bicarbonate/water (50 mL), brine (50 mL), dried (magnesium sulfate), concentrated in vacuo and chromatographed (silica gel,

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ethyl acetate/hexane, 1:9) to afford the title compound (1.0g, 23%) as a clear liquid;

^1H NMR (CDCl_3): δ 1.05 (6H, s), 1.31 (3H, s), 1.52-1.62 (4H, m), 3.88-3.98 (4H, m), 9.45 (1H, s).

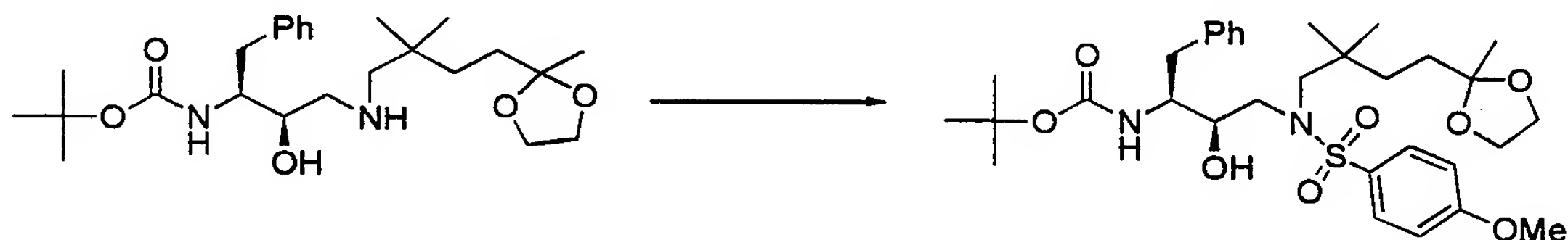
5 Step 2:



tert-Butyl N-((1S,2R)-1-benzyl-3-[2,2-dimethyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl]amino-2-hydroxypropyl) carbamate

The product from Step 1 was subjected to Step 2
 10 (Example (Compound 201)) to afford the title compound as
 a thick oil; ^1H NMR ($\text{DMSO}-d_6$): δ 0.80 (6H, s), 1.17-1.27
 (14H, m), 1.38 (1H, br), 1.42-1.50 (2H, m), 2.22 (2H, s),
 2.45-2.59 (3H, m), 2.96 (1H, dd), 3.36-3.55 (2H, m),
 3.75-3.85 (4H, m), 4.76 (1H, br s), 6.71 (1H, d), 7.08-
 15 7.24 (5H, m) ; MS 451 (MH+)

Step 3:

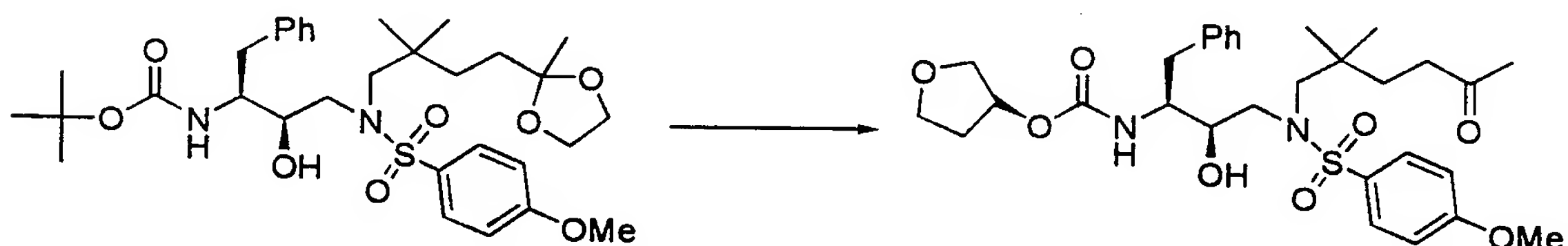


tert-Butyl N-((1S,2R)-1-benzyl-3-[2,2-dimethyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl][(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

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The product from Step 2 was subjected to Step 3 (Example (Compound 201)) to give the title compound as a white solid; ^1H NMR ($\text{DMSO}-d_6$): δ 0.88 (6H, s), 1.19 (3H, s), 1.22 (9H, s), 1.28 (2H, t), 1.44-1.56 (2H, m), 2.42-2.49 (1H, m), 2.77 (1H, d), 2.87-2.98 (2H, m), 3.25-3.40 (3H, m), 3.65-3.73 (1H, m), 3.82 (7H, s), 4.95 (1H, d), 6.60 (1H, d), 7.03-7.23 (7H, m), 7.77 (2H, d) ; MS 621 (MH $^+$) .

10 Step 4:



(3S)-Tetrahydro-3-furanylN-((1S,2R)-1-benzyl-3-(2,2-dimethyl-5-oxohexyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

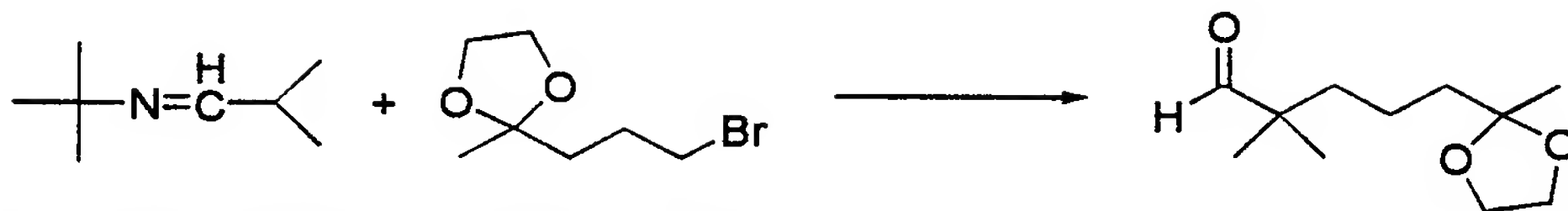
15 The product from Step 3 was subjected to the procedure used in Example (Compound 205) to provide the title compound as a white foam;

^1H NMR ($\text{DMSO}-d_6$): δ 0.87 (3H, s), 0.9 (3H, s), 1.47 (2H, t), 1.75 (1H, dt), 2.03-2.08 (1H, m), 2.1 (3H, s), 2.35-2.45 (3H, m), 2.75 (1H, d), 2.85-2.95 (2H, m), 3.3-3.5 (3H, m), 3.57 (1H, dd), 3.64 (1H, dd), 3.66 (1H, dd), 3.70 (1H, dd), 3.75 (1H, dd), 3.82 (3H, s), 4.9 (1H, brt), 5.1 (1H, d), 7.08 (2H, d), 7.10-7.25 (6H, m), 7.7 (2H, d); MS: 591.1 (MH $^+$) .

Example (Compound 209)

25 Step 1:

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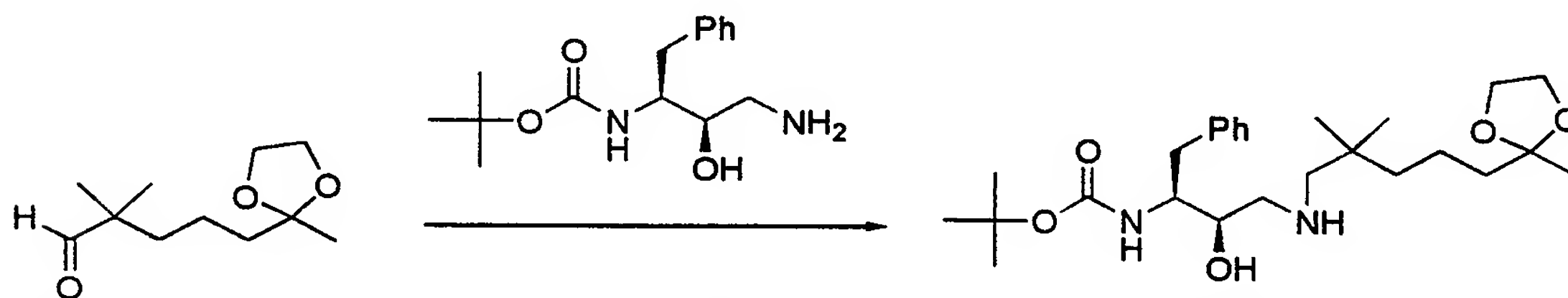


2,2-Dimethyl-5-(2-methyl-1,3-dioxolan-2-yl)pentanal

The title compound was prepared by procedure used in Example (Compound 202) using 2-(3-chloropropyl)-2-methyl-1,3-dioxolane as alkylating agent.

- 5 ^1H NMR (CDCl_3): δ 1.05 (6H, s), 1.24-1.36 (5H, m), 1.44-1.50 (2H, m), 1.60-1.65 (2H, m), 3.88-3.98 (4H, m), 9.45 (1H, s)

Step 2:



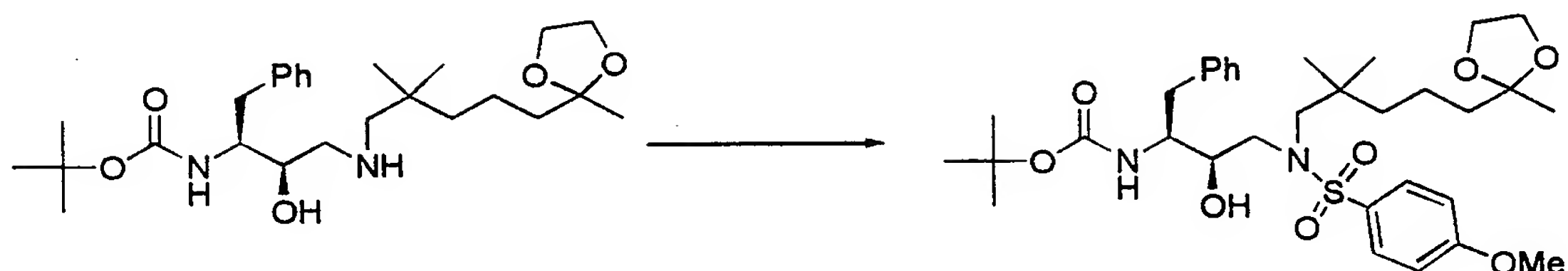
- 10 **tert-Butyl N-((1S,2R)-1-benzyl-3-[2,2-dimethyl-5-(2-methyl-1,3-dioxolan-2-yl)pentyl]amino-2-hydroxypropyl)carbamate**

The product from step 1 was subjected to Step 2 (Example (Compound 201)) to provide the title compound as a thick oil; ^1H NMR ($\text{DMSO}-d_6$): δ 0.80 (6H, s), 1.10-1.30 (16H, m), 1.42 (1H, brs), 1.45-1.52 (2H, m), 2.22 (2H, s), 2.47-2.60 (3H, m), 2.95 (1H, dd), 3.37-3.57 (2H, m), 3.75-3.85 (4H, m), 4.76 (1H, br s), 6.75 (1H, dd), 7.08-7.26 (5H, m) ; MS: 465 (MH^+).

15

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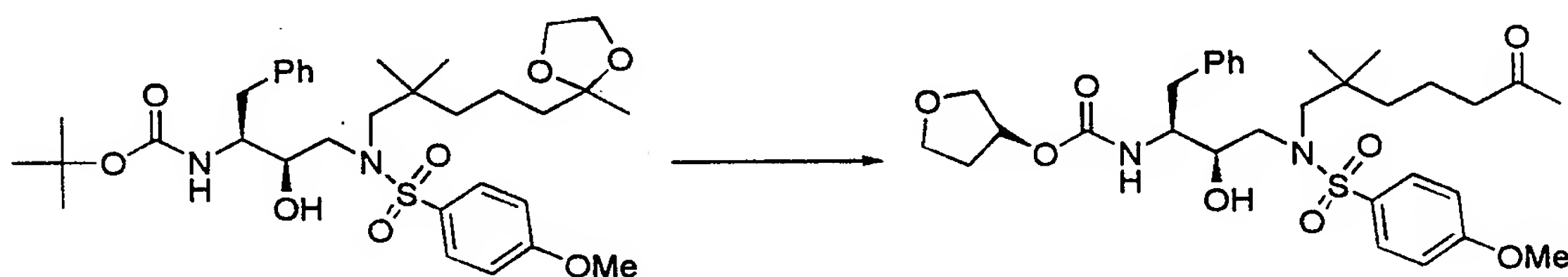
Step 3:



The product from the previous step was subjected to Step 3 (Example (Compound 201)) to afford the title compound as a white solid;

- 5 ^1H NMR ($\text{DMSO}-d_6$): δ 0.88 (6H, s), 1.14-1.30 (16H, m), 1.44-1.51 (2H, m), 2.42-2.48 (1H, m), 2.77 (1H, d), 2.87-2.98 (2H, m), 3.25-3.40 (3H, m), 3.65-3.75 (1H, m), 3.75-3.85 (7H, m), 4.96 (1H, d), 6.60 (1H, dd), 7.04-7.24 (7H, m), 7.72 (2H, d); MS: 635 (MH^+).

10 Step 4:



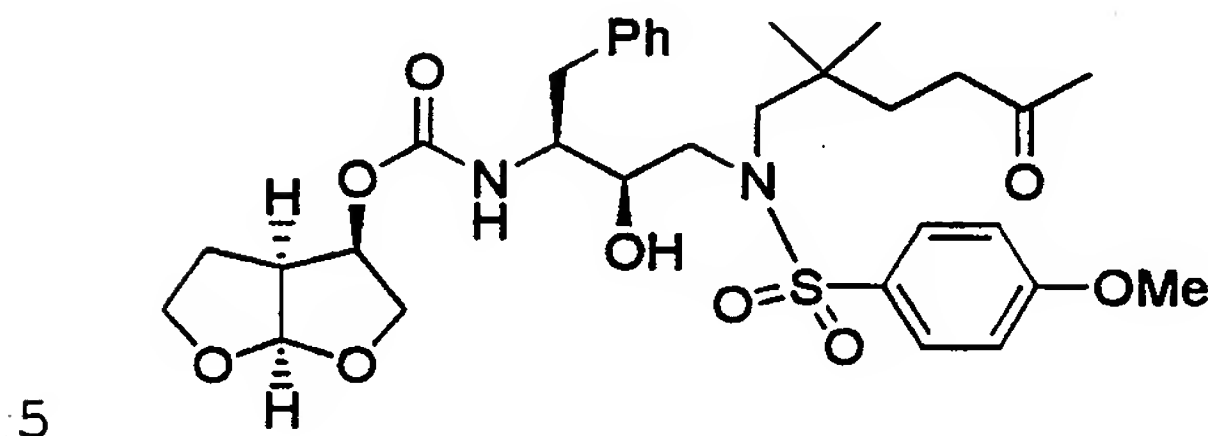
The product from the previous step was subjected to the Step used in Example (Compound 205) to give the title compound as a thick oil;

- 15 ^1H NMR ($\text{DMSO}-d_6$): δ 0.89 (3H, s), 1.15 (2H, dd), 1.3-1.5 (2H, m), 1.75 (1H, dd), 1.95-2.05 (1H, m), 2.08 (3H, s), 2.35 (2H, t), 2.4-2.5 (1H, m), 2.75 (1H, d), 2.93 (2H, brd), 3.30-3.45 (4H, m), 3.6 (1H, dd), 3.63 (1H, dd), 3.7

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(1H, d), 3.72-3.77 (1H, m), 3.81 (3H, s), 4.9 (1H, brt), 5.0 (1H, d), 7.08 (2H, d), 7.10-7.25 (6H, m), 7.7 (2H, d); MS: 604.9 (MH⁺)

Example (Compound 210)



(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(2,2-dimethyl-5-oxohexyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

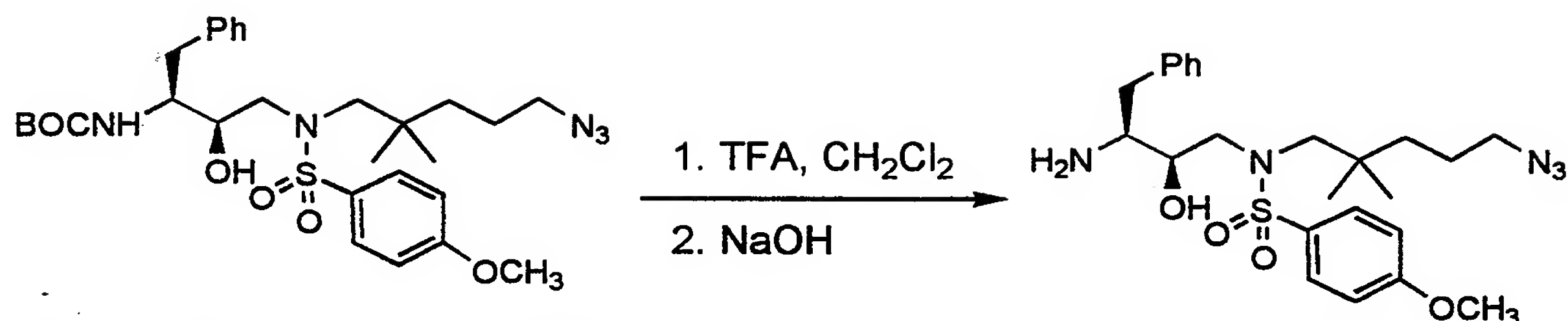
The product from Step 3 (Example (Compound 208)) was subjected to a procedure similar to that of Example (Compound 205) (except for the use of (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-p-nitrophenyl carbonate in the acylation step) to give the title compound as a foam;

¹H NMR (DMSO-d₆): δ 0.82 (3H, s), 0.86 (3H, s), 1.2 (1H, dd), 1.3-1.4 (1H, m), 1.43 (2H, t), 2.04 (3H, s), 2.3-2.5 (3H, m), 2.65-2.80 (2H, m), 2.89 (1H, dd), 2.92 (1H, dd), 3.3-3.4 (3H, m), 3.42-3.50 (1H, m), 3.57 (2H, dd), 3.73 (2H, dd), 3.9 (3H, s), 4.8 (1H, quartet), 5.1 (1H, d), 5.48 (1H, d), 7.05 (2H, d), 7.10-7.25 (6H, m), 7.7 (2H, d); MS: 633 (MH⁺).

Example (Compound 211)

Step 1:

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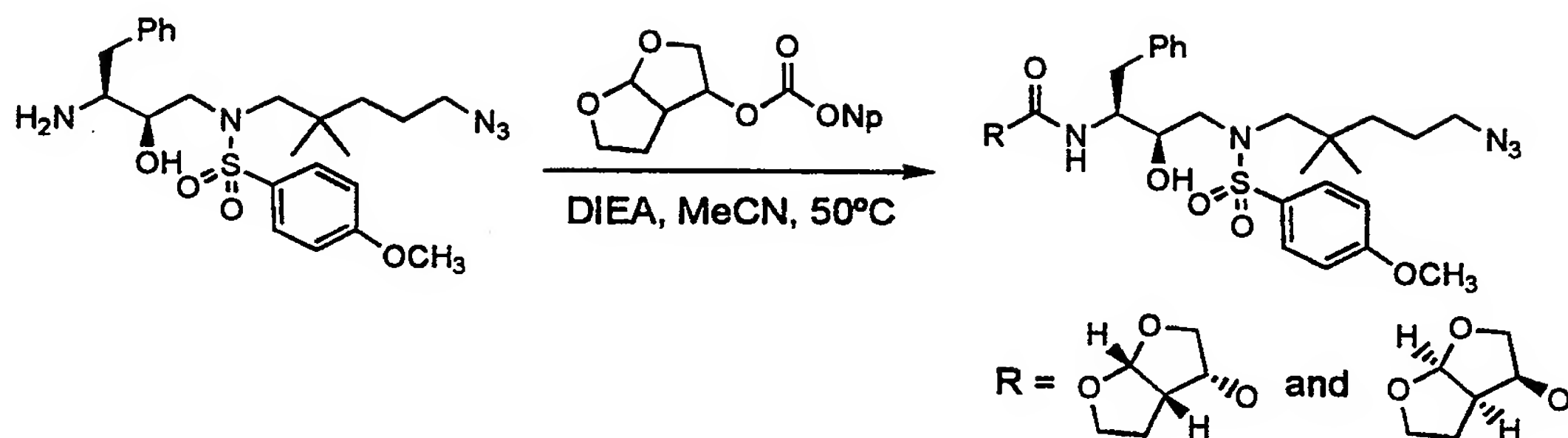
N-[(2*R*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl]-*N*-(5-azido-2,2-dimethylpentyl)-4-methoxybenzenesulfonamide

A solution of 0.49 g (1.7 mmol) of *tert*-butyl *N*-((1*S*,2*R*)-3-(5-azido-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate (Example (Compound 203)) in 10 mL of CH₂Cl₂ was treated with 10 mL of TFA and the solution was stirred at RT. After 3 hours the solution was concentrated in vacuo. The residue was dissolved in CH₂Cl₂. The solution was washed with 1M aqueous NaOH (1x), water (2x), dried over MgSO₄, and concentrated in vacuo. The crude material was subjected to flash chromatography (SiO₂, 97:3 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.32 g of the desired compound as white foam.

¹H NMR (DMSO-*d*₆): δ 7.66 (2H), 7.22 (2H), 7.15 (3H), 7.03 (2H), 4.64 (1H), 3.79 (3H), 3.62-3.20 (5H), 3.03 (1H), 2.85 (1H), 2.74-2.60 (2H), 2.27 (1H), 1.48 (2H), 1.23 (2H), 1.10 (2H), 0.84 (6H). MS (ESI): 490 (M+H).

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Step 2:



(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-azido-2,2-dimethylpentyl) [(4-methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate and

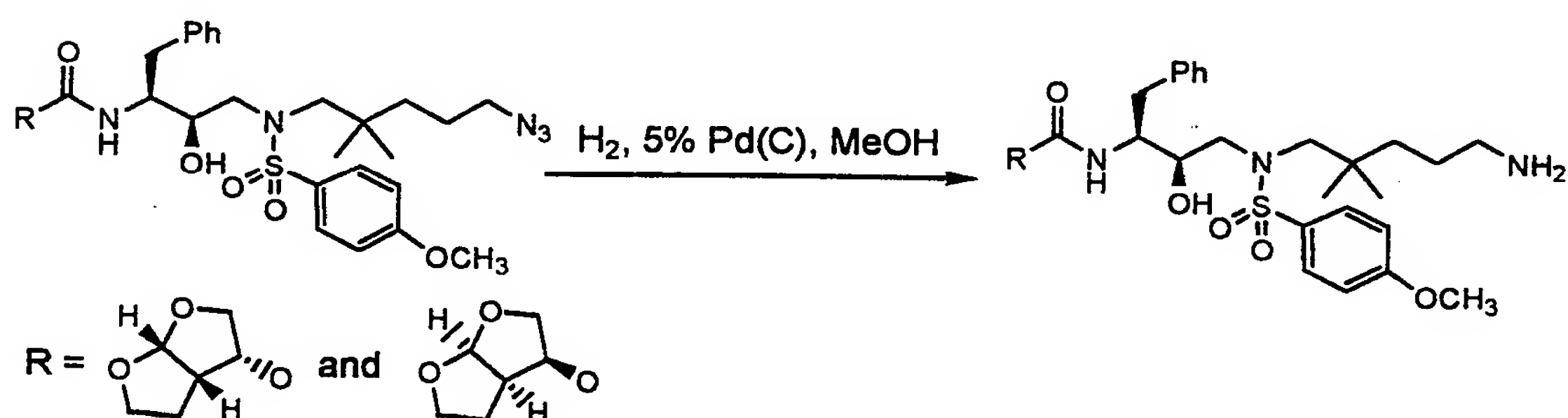
5 (3S,3aR,6aS) hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-azido-2,2-dimethylpentyl) [(4-methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate

A solution of 0.31 g (0.63 mmol) of N-[(2R,3S)-3-amino-2-hydroxy-4-phenylbutyl]-N-(5-azido-2,2-dimethylpentyl)-4-methoxybenzenesulfonamide (step 1),
 10 0.20 g (0.67 mmol) of a 1:1 mixture of (3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl (4-nitrophenyl) carbonate and (3S,3aR,6aS) hexahydrofuro[2,3-b]furan-3-yl (4-nitrophenyl) carbonate, and 0.18 mL of N,N-diisopropylethylamine in 10 mL of acetonitrile was
 15 stirred at 50°C under nitrogen. After 24 hours the solution was cooled to RT and concentrated in vacuo. The residue was dissolved in CH₂Cl₂. The solution was washed with 0.2M aqueous NaOH (4x), water (2x), dried over MgSO₄,
 20 and concentrated to dryness. The crude product was subjected to flash chromatography (SiO₂, 1:1 hexane/EtOAc) to afford 0.27 g (66%) of the desired product as a white foam.

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^1H NMR (CDCl_3): δ 7.70 (2H), 7.32-7.11 (5H), 7.00 (2H), 5.62 (1H), 4.97 (2H), 4.18-3.97 (2H), 3.96-3.72 (6H), 3.70-3.42 (2H), 3.30-3.01 (4H), 3.00-2.63 (4H), 1.99-1.72 (1H), 1.53 (2H), 1.41-1.15 (3H), 0.90 (6H). LCMS (ESI):

5 646 (M+H).

Example (Compound 212)

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and

(3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate

15 A solution of 0.25 g (0.39 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-azido-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and

(3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-azido-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate (Example

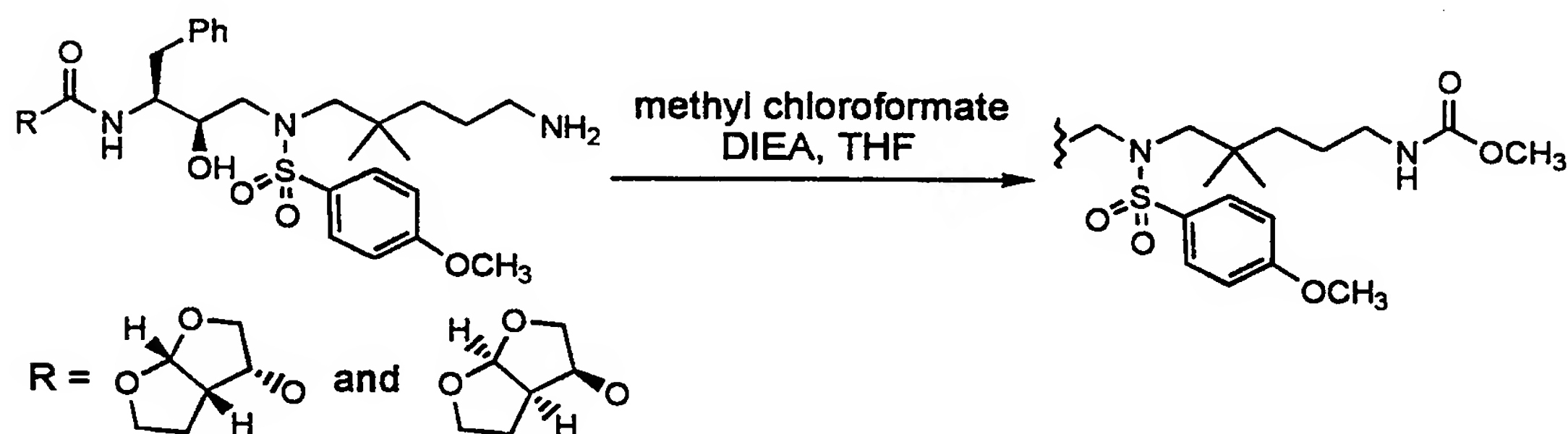
20 (Compound 211)) in 40 mL of MeOH under nitrogen was treated with 50 mg of 5% Pd(C). The resulting mixture was subjected to hydrogenation at 30 psi. After 2 hours

-176-

the vessel was purged with nitrogen, catalyst removed by filtration through celite and the filtrate concentrated in vacuo to afford 0.23 g (96%) of the desired product as a light yellow foam.

5 ^1H NMR (CDCl_3): δ 7.75 (2H), 7.38-7.12 (5H), 7.03 (2H), 5.65 (1H), 5.02 (1H), 4.13-3.44 (8H), 3.38-2.55 (10H), 2.02-1.08 (6H), 1.07-0.83 (6H). LCMS (ESI): 620 (M+H).

Example (Compound 213)



(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-2-hydroxy-3-5-[(methoxycarbonyl)amino]-2,2-dimethylpentyl[(4-methoxyphenyl)sulfonyl]aminopropyl)carbamate and (3S,3aR,6aS) hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-2-hydroxy-3-5-[(methoxycarbonyl)amino]-2,2-dimethylpentyl[(4-methoxyphenyl)sulfonyl]aminopropyl)carbamate

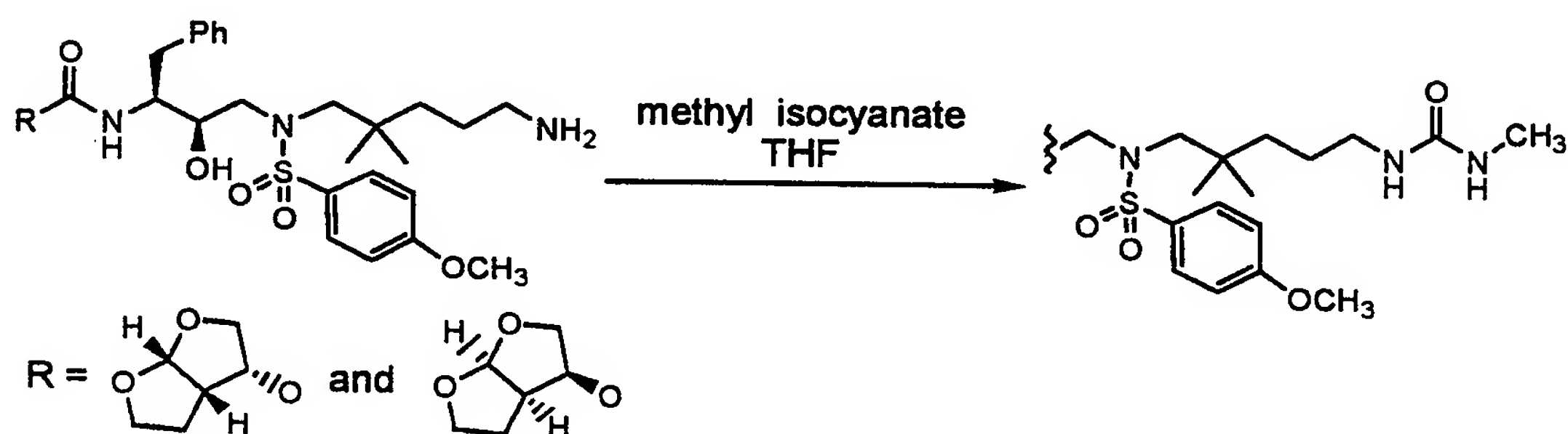
A solution of 20 mg (0.032 mmol) of a 1:1 mixture of (3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3S,3aR,6aS) hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate (Example (Compound 212)) in 1.5 mL of anhydrous THF at 0°C was

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treated with 6.3 μL (0.036 mmol) of *N,N*-diisopropylethylamine followed by 2.8 μL (0.036 mmol) of methyl chloroformate. The resulting solution was allowed to warm to RT with stirring. After 2.5 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO_2 , 97:3 $\text{CH}_2\text{Cl}_2/\text{MeOH}$) to afford 16 mg (74%) of the desired product as a white foam.

^1H NMR (CDCl_3): δ 7.70 (2H), 7.29–7.12 (5H), 6.99 (2H), 5.61 (1H), 5.46–5.21 (1H), 5.10–4.88 (2H), 4.20–3.42 (12H), 3.23–2.61 (9H), 1.99–1.13 (6H), 0.91 (3H), 0.83 (3H). LCMS (ESI): 678 (M+H).

Example (Compound 214)



(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-5-[(methylamino)carbonyl]aminopentyl) [(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate and (3*S*,3*aR*,6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-5-[(methylamino)carbonyl]aminopentyl) [(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

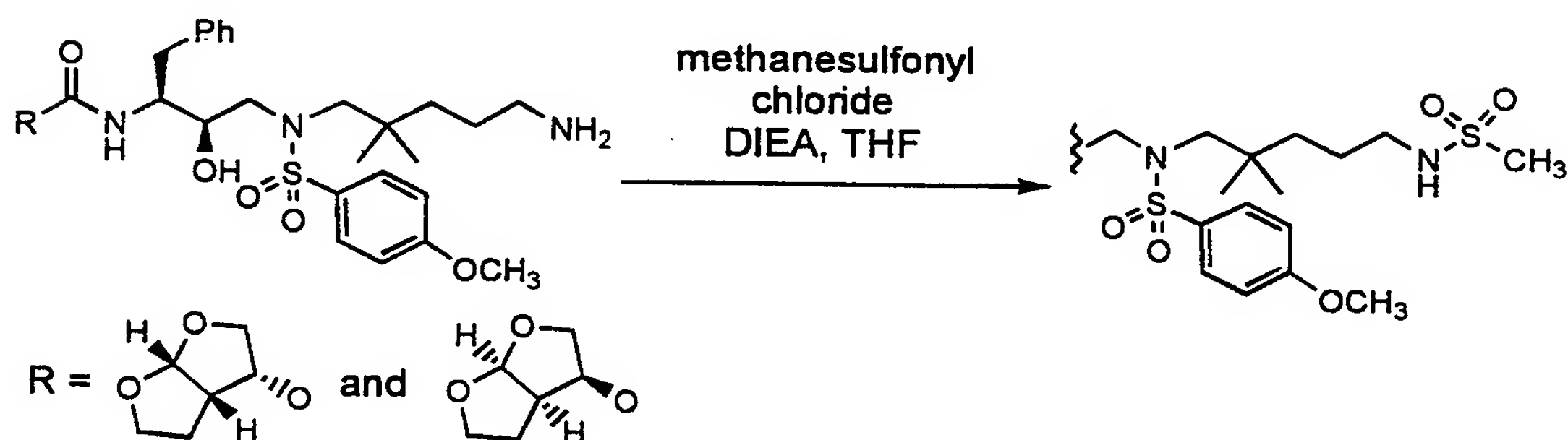
A solution of 20 mg (0.032 mmol) of a 1:1 mixture of (3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl) [(4-

-178-

methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl)
 carbamate and (3*S*,3*aR*,6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl
N-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl) [(4-
 methoxyphenyl) sulfonyl] amino-1-benzyl-2-
 5 hydroxypropyl) carbamate (Example (Compound 212)) in 1.5
 mL of anhydrous THF at 0°C was treated with 2.0 µL (0.032
 mmol) of methyl isocyanate. The resulting solution was
 allowed to warm to RT with stirring. After 2.5 hours the
 solution was concentrated *in vacuo* and the residue
 10 subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH)
 to afford 15 mg (69%) of the desired product as a white
 foam.

¹H NMR (CDCl₃): δ 7.70 (2H), 7.30-7.10 (5H), 7.00 (2H),
 5.83-5.48 (2H), 5.10-4.50 (2H), 4.30-3.43 (10H), 3.30-
 15 2.62 (12H), 2.10-1.18 (6H), 0.93 (3H), 0.84 (3H).
 LCMS (ESI): 677 (M+H).

Example (Compound 215)



(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-
 benzyl-3-2,2-dimethyl-5-[(methanesulfonyl) amino]pentyl [(4-
 20 methoxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate
 and (3*S*,3*aR*,6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-
 ((1*S*,2*R*)-1-benzyl-3-2,2-dimethyl-5-[(methanesulfonyl)

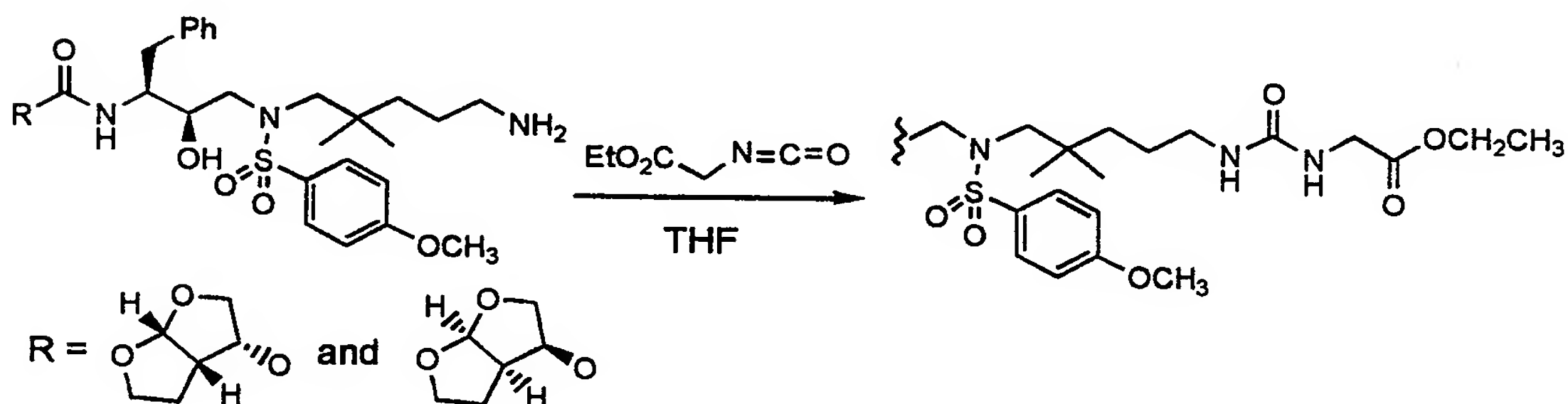
-179-

amino]pentyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

A solution of 20 mg (0.032 mmol) of a 1:1 mixture of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate (Example 10 (Compound 212)) in 1.5 mL of anhydrous THF at 0°C was treated with 6.0 µL (0.032 mmol) of *N,N*-diisopropylethylamine followed by 2.5 µL (0.032 mmol) of methanesulfonyl chloride. The resulting solution was allowed to warm to RT with stirring. After 2.5 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 97:3 CH₂Cl₂/MeOH) to afford 15 mg (68%) of the desired product as a white foam.

¹H NMR (CDCl₃): 7.69 (2H), 7.31-7.10 (5H), 7.00 (2H), 5.61 (1H), 5.18-5.02 (1H), 4.92 (2H), 4.07 (1H), 3.99-3.34 (8H), 3.21-2.63 (12H), 2.04-1.20 (6H), 0.93 (3H), 0.87 (3H). LCMS (ESI): 698 (M+H).

Example (Compound 216)



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Ethyl (3*S*,4*R*)-1-[(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl)sulfonyl]-8,8-dimethyl-1,13-dioxo-2,6,12,14-tetraazahexadecan-16-oate and ethyl (3*S*,4*R*)-1-[(3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl)sulfonyl]-8,8-dimethyl-1,13-dioxo-2,6,12,14-tetraazahexadecan-16-oate

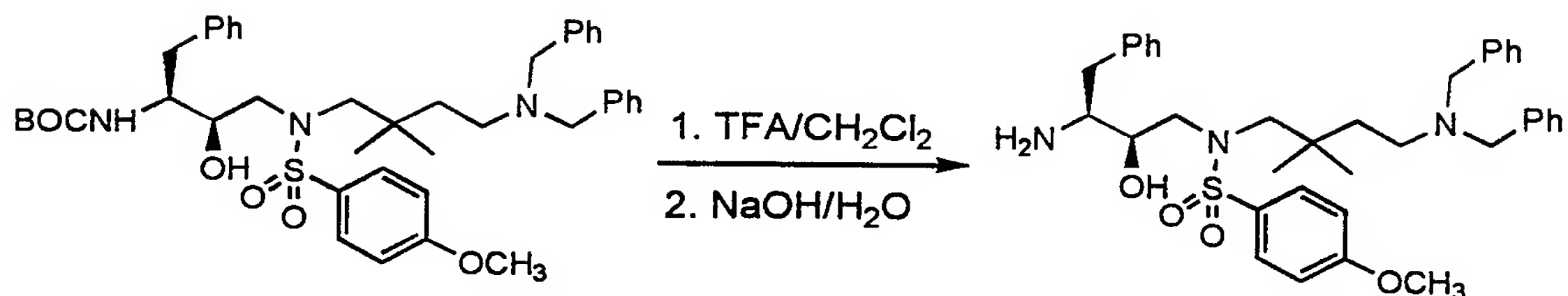
A solution of 33 mg (0.053 mmol) of a 1:1 mixture of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate (Example 15 (Compound 212)) in 1.5 mL of anhydrous THF was treated with 6.5 μ L (0.058 mmol) of ethyl isocyanato acetate and the resulting solution was stirred at RT. After 18 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 97:3 CH₂Cl₂/2M NH₃ in MeOH) to afford 31 mg (78%) of the desired compound as a white foam.

¹H NMR (CDCl₃): 7.76 (2H), 7.35-7.16 (5H), 7.04 (2H), 5.80-5.42 (2H), 5.02 (1H), 4.37-3.46 (13H), 3.40-2.63 (9H), 2.18-1.78 (2H), 1.70-1.22 (7H), 1.01 (3H), 0.92 (3H). LCMS (ESI): 749 (M+H).

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Example (Compound 217)

Step 1:



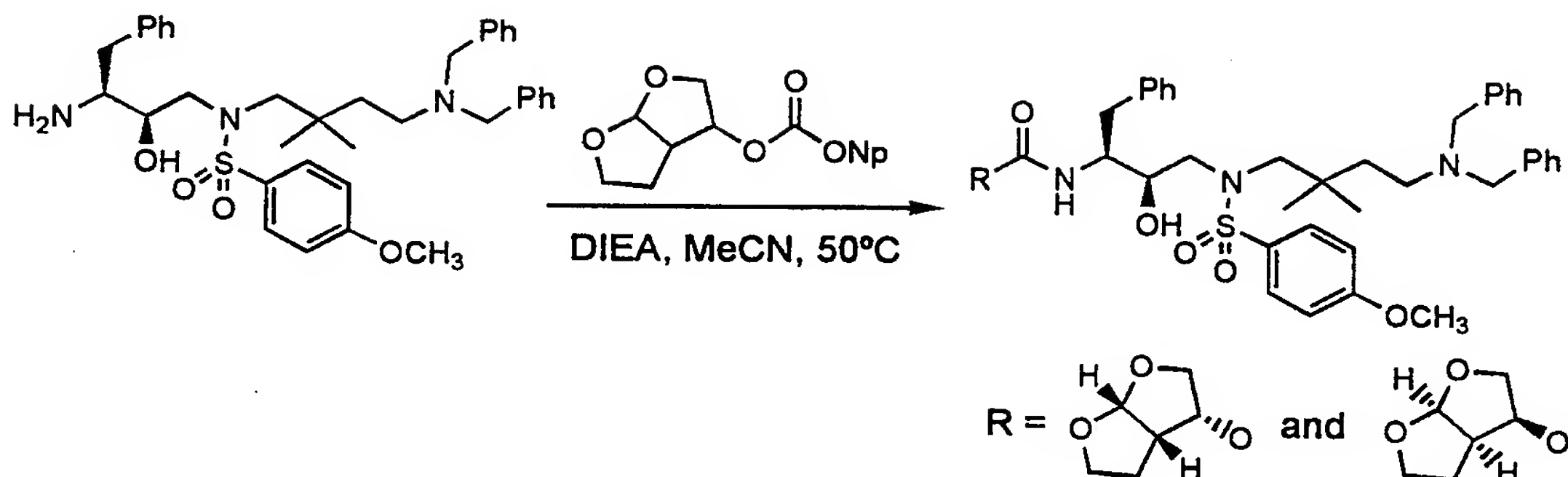
5 *N*-[(2*R*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl]-*N*-[4-(dibenzylamino)-2,2-dimethylbutyl]-4-methoxybenzenesulfonamide

A solution of 1.00 g (1.37 mmol) of *tert*-butyl *N*-((1*S*,2*R*)-1-benzyl-3-[4-(dibenzylamino)-2,2-dimethylbutyl][(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate in 40 mL of 1:1 CH₂Cl₂/TFA was stirred at RT for 2 hours. The solution was then concentrated to dryness and the residue dissolved in CH₂Cl₂. The resulting solution was washed with 1 M aqueous NaOH (1x), water (3x), dried over MgSO₄ and concentrated *in vacuo* to afford 0.81 g (94%) of the desired amine as a thick oil.

¹H NMR (CDCl₃): 7.76 (2H), 7.44-7.19 (15H), 7.01 (2H), 4.00 (1H), 3.92 (3H), 3.63 (4H), 3.34-2.88 (6H), 2.53 (3H), 1.64 (2H), 0.92 (6H).

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Step 2:



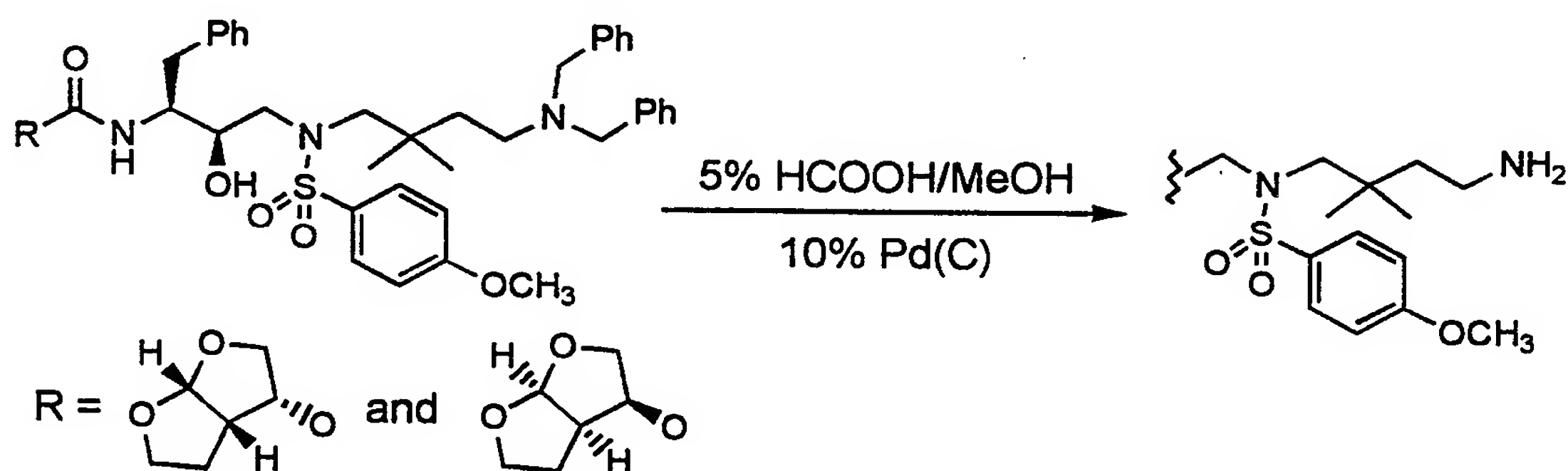
(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-[4-(dibenzylamino)-2,2-dimethylbutyl]-(4-methoxyphenyl)sulfonyl)amino-2-hydroxypropyl) carbamate
 5 and (3S,3aR,6aS) hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-[4-(dibenzylamino)-2,2-dimethylbutyl]-(4-methoxyphenyl)sulfonyl)amino-2-hydroxypropyl) carbamate

A solution of 0.81 g (1.3 mmol) of N-[(2R,3S)-3-amino-2-hydroxy-4-phenylbutyl]-N-[4-(dibenzylamino)-2,2-dimethylbutyl]-4-methoxybenzenesulfonamide (step 1),
 10 0.40 g (1.4 mmol) of a 1:1 mixture of (3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl (4-nitrophenyl)carbonate and (3S,3aR,6aS) hexahydrofuro[2,3-b]furan-3-yl (4-nitrophenyl)carbonate, and 0.37 mL (2.1 mmol) of N,N-diisopropylethylamine in 20 mL of acetonitrile was
 15 stirred at 50°C. After 3.5 hours the solution was cooled to RT and concentrated to dryness. The residue was dissolved in CH₂Cl₂ and the solution washed with 1 M aqueous NaOH (4X), water (3x), dried over MgSO₄ and
 20 concentrated *in vacuo*. The crude product was subjected to flash chromatography (SiO₂, 1:1 hexane/EtOAc) to afford 0.85 g (84%) of the desired product as a white foam.

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^1H NMR (CDCl_3): δ 7.71 (2H), 7.58–7.13 (15H), 7.01 (2H), 5.66 (1H), 5.07–4.60 (2H), 4.40–3.40 (12H), 3.27–2.63 (9H), 2.46 (1H), 2.20–1.45 (4H), 0.87 (6H). LCMS (ESI): 786 (M+H).

5 Step 3:



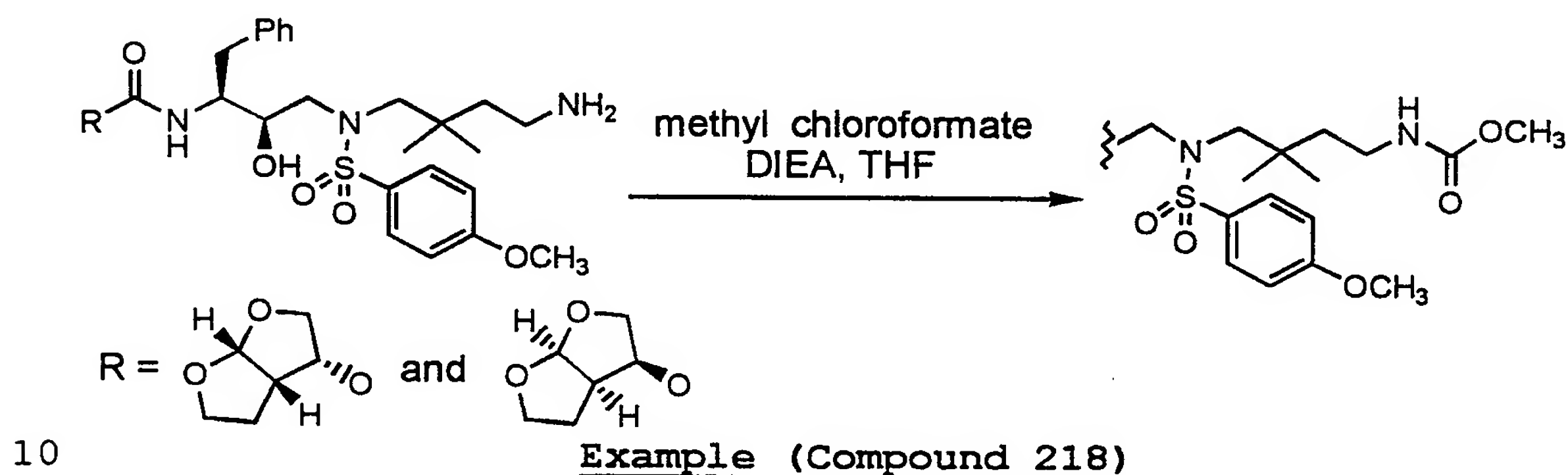
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and
 (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate

A solution of 0.78 g (0.99 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-[4-(dibenzylamino)-2,2-dimethylbutyl][(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-[4-(dibenzylamino)-2,2-dimethylbutyl][(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate (Step 2) in 50 mL of 5% formic acid/MeOH under nitrogen was treated with 1.00 g of 10% Pd(C) and the resulting mixture was stirred at RT. After 18 hours the catalyst was removed by filtration through celite and the filtrate concentrated

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to dryness in vacuo. The residue was dissolved in CH₂Cl₂ and the solution was washed with 1M aqueous NaOH (1x), water (2X), dried over MgSO₄, and concentrated to dryness. The crude product was subjected to flash chromatography
 5 (SiO₂, 9:1 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.45 g (75%) of the desired product as a white foam.

¹H NMR (CDCl₃): δ 7.70 (2H), 7.21 (5H), 6.99 (2H), 5.60 (1H), 5.00 (1H), 4.18-3.40 (8H), 3.30-2.63 (10H), 2.02-1.30 (4H), 1.01 (3H), 0.88 (3H). MS(ESI): 606 (M+H).



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-2-hydroxy-3-4-[(methoxycarbonyl)amino]-2,2-dimethylbutyl[(4-methoxyphenyl)sulfonyl]aminopropyl) carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl
 15 N-((1S,2R)-1-benzyl-2-hydroxy-3-4-[(methoxycarbonyl)amino]-2,2-dimethylbutyl[(4-methoxyphenyl)sulfonyl]aminopropyl) carbamate

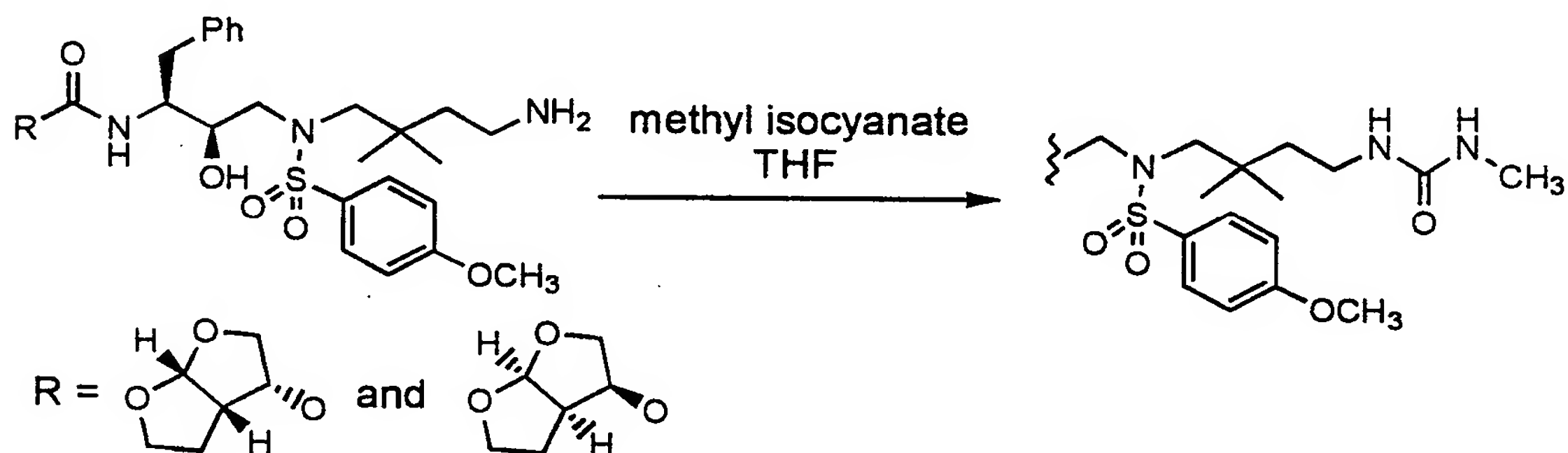
A solution of 25 mg (0.041 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-
 20 ((1S,2R)-3-(4-amino-2,2-dimethylbutyl) [(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-

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(4-amino-2,2-dimethylbutyl) [(4-methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate (Example (Compound 217)) in 1.5 mL of anhydrous THF was cooled to 0°C and was treated with 8.0 µL (0.045 mmol) of *N,N*-diisopropylethylamine followed by 3.5 µL (0.045 mmol) of methyl chloroformate. The solution was allowed to warm to RT with stirring. After 1.5 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 18 mg (67%) of the desired product as a white foam.

¹H NMR (CDCl₃): 7.75 (2H), 7.37-7.18 (5H), 7.03 (2H), 5.65 (1H), 5.39-4.90 (3H), 4.20-3.50 (12H), 3.38-2.69 (9H), 2.18-1.39 (4H), 1.09 (3H), 0.95 (3H). MS (ESI): 664 (M+H).

15

Example (Compound 219)

(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-4-[(methylamino)carbonyl]aminobutyl) [(4-methoxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate and
 (3*S*,3*aR*,6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-4-[(methylamino)carbonyl]

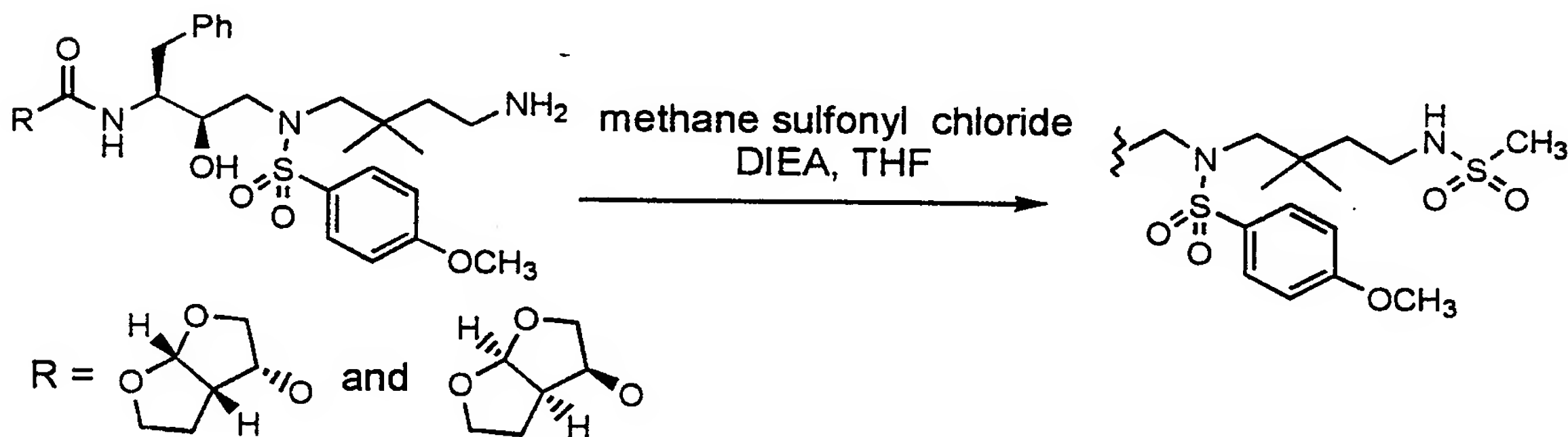
-186-

aminobutyl) [(4-methoxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate

A solution of 25 mg (0.041 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-dimethylbutyl) [(4-methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-dimethylbutyl) [(4-methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate (Example 10 (Compound 217)) in 1.5 mL of anhydrous THF was cooled to 0°C and was treated with 2.5 µL (0.043 mmol) of methyl isocyanate. The resulting solution was allowed to warm to RT with stirring. After 2 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH) to afford 16 mg (59%) of the desired product as a white foam.

¹H NMR (CDCl₃): δ 7.74 (2H), 7.37-7.14 (5H), 7.02 (2H), 5.75-5.50 (2H), 5.01 (1H), 4.19-3.49 (9H), 3.40-2.70 (12H), 2.18-1.23 (4H), 1.03 (3H), 0.97 (3H). MS (ESI): 663 (M+H).

Example (Compound 220)



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-2,2-dimethyl-4-[(methylsulfonyl) amino]butyl [(4-

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methoxyphenyl) sulfonyl]amino-2-hydroxypropyl) carbamate
 and (3*S*, 3*aR*, 6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-
 ((1*S*, 2*R*)-1-benzyl-3-2,2-dimethyl-4-[(methanesulfonyl)
 amino]butyl[(4-methoxyphenyl) sulfonyl]amino-2-
 5 hydroxypropyl) carbamate

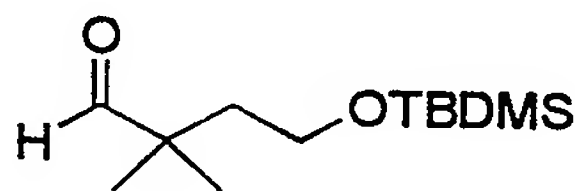
A solution of 25 mg (0.041 mmol) of a 1:1
 mixture of (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-
 ((1*S*, 2*R*)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)
 sulfonyl]amino-1-benzyl-2-hydroxypropyl) carbamate and
 10 (3*S*, 3*aR*, 6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-3-
 (4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl) sulfonyl]
 amino-1-benzyl-2-hydroxypropyl) carbamate (Example
 (Compound 217)) in 1.5 mL of anhydrous THF was cooled to
 0°C and was treated with 7.0 µL (0.041 mmol) of *N,N*-
 15 diisopropylethylamine followed by 3.2 µL (0.041 mmol) of
 methane sulfonyl chloride. The solution was allowed to
 warm to RT with stirring. After 2 hours the solution was
 concentrated *in vacuo* and the residue was subjected to
 flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH) to afford 20
 20 mg (71%) of the desired product as a white foam.

¹H NMR (CDCl₃): δ 7.76 (2H), 7.38-7.19 (5H), 7.06 (2H),
 5.67 (1H), 5.29-4.99 (3H), 4.31-3.54 (10H), 3.32-2.63
 (11H), 2.10-1.37 (4H), 1.04 (3H), 0.99 (3H). MS (ESI):
 684 (M+H).

25

Example (Compound 221)

Step 1:



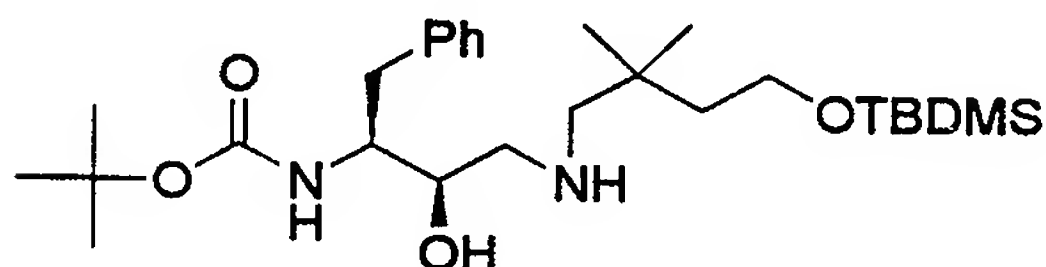
4-*tert*-Butyldimethylsilyloxy-2,2-dimethylbutanal

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The title compound was prepared by a procedure similar to Example (Compound 202) using 2-*tert*-butyldimethyl silyloxybromoethane as alkylating agent.

¹H NMR (CDCl₃): δ -0.02 (6H, s), 0.84 (9H, s), 1.04 (6H, s), 1.74 (2H, t), 3.61 (2H, t), 9.43 (1H, s)

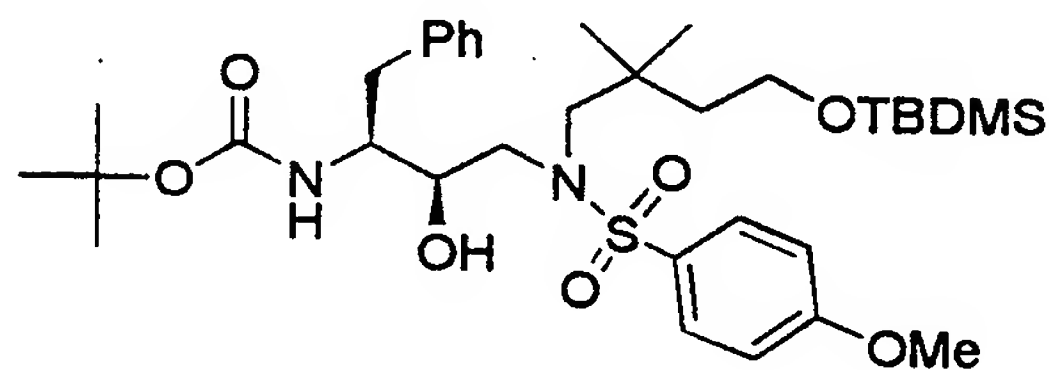
Step 2:



The product from Step 1 was subjected to Step 2 (Example (Compound 201)) to give the title compound as a foam;

¹H NMR (DMSO-*d*₆): δ 0.01 (6H, s), 0.82 (15H, s), 1.22 (9H, s), 1.42 (2H, t), 2.23 (2H, s), 2.43-2.60 (3H, m), 2.95 (1H, dd), 3.33-3.58 (2H, m), 3.60 (2H, t), 4.72 (1H, br s), 6.71 (1H, d), 7.07-7.25 (5H, m) ; MS: 495 (MH⁺).

Step 3:



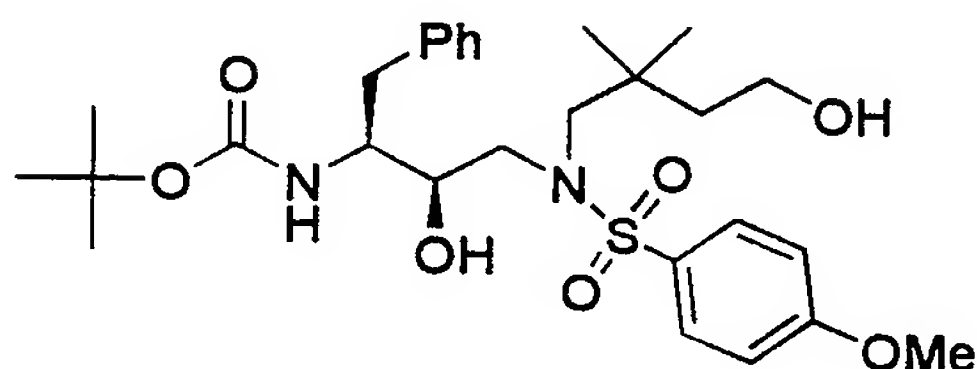
tert-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-(4-*tert*-butyldimethylsilyloxy-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

The product from Step 2 was subjected to Step 3 (Example (Compound 201)) to provide the title compound as a foam;

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¹H NMR (DMSO-*d*₆): δ 0.00 (6H, s), 0.82 (9H, s), 0.93 (6H, s), 1.20 (9H, s), 1.48 (2H, t), 2.41-2.49 (1H, m), 2.74-2.96 (3H, m), 3.24-3.40 (3H, m), 3.63 (2H, t), 3.67-3.75 (1H, m), 3.81 (3H, s), 4.95 (1H, broad), 6.61 (1H, d),
5 7.04 (2H, d), 7.08-7.24 (2H, d), 7.71 (2H, d) ; MS: 665 (MH⁺) .

Step 4:



tert-Butyl N-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-4-hydroxybutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate
10

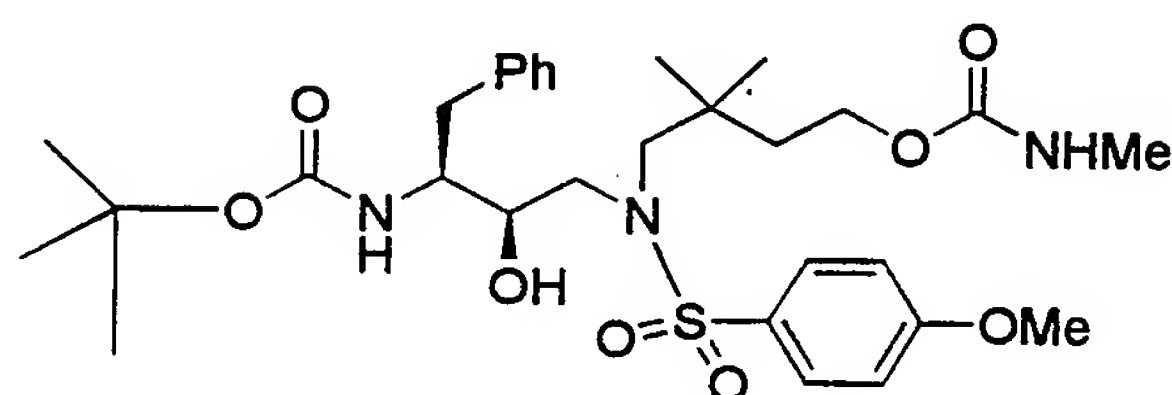
To a stirred solution of the product of Step 3 (3.47g, 5.2mmol) in tetrahydrofuran (25 mL) was added tetrabutylammonium fluoride (6.2 mL, 1M in tetrahydrofuran) over 15 minutes. After stirring at
15 ambient temperature for 2 hours, the reaction was concentrated *in vacuo* and partitioned in ether/water (1:1; 200 mL). The layers were separated and the aqueous phase was extracted with ether (50 mL). The combined organic layers were washed with water (100 mL), dried
20 (magnesium sulfate), concentrated *in vacuo* and purified by silica gel chromatography (4% methanol in chloroform) to afford the title compound (2.78g, 97%) as a white foam.

¹H NMR (DMSO-*d*₆): δ 0.98 (6H, s), 1.27 (9H, s), 1.51 (2H, t), 2.46-2.57 (1H, m), 2.81-3.06 (3H, m), 3.32-3.55 (5H, m), 3.71-3.82 (1H, m), 3.87 (3H, s), 4.32 (1H, t), 5.01

-190-

(1H, d), 6.66 (1H, d), 7.12 (2H, d), 7.16-7.31 (5H, m), 7.77 (2H, d) ; MS: 551 (MH⁺).

Step 5:



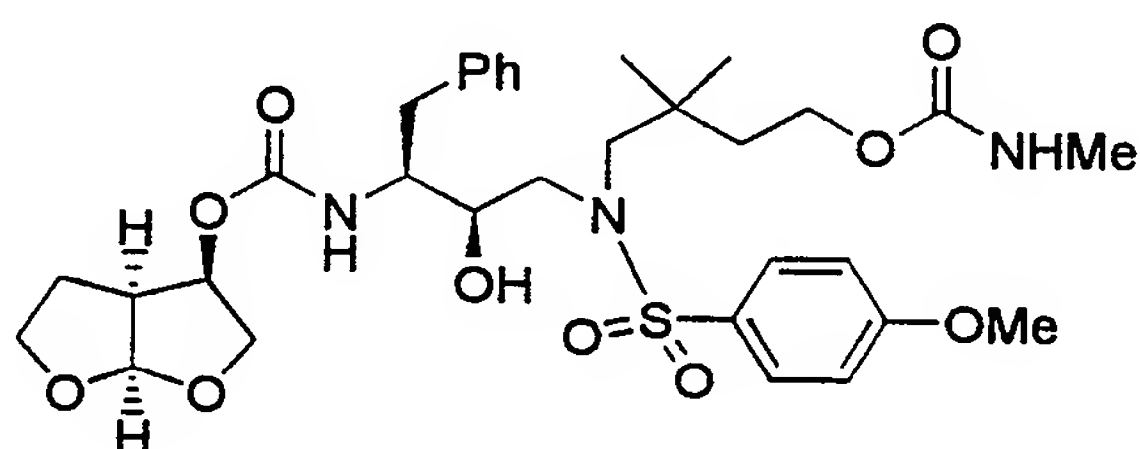
5 *tert*-Butyl *N*-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-4-*N'*-methylcarbamoyloxybutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

To a solution of the product from Step 4 (500 mg, 0.9 mmol) in dichloromethane (4 mL) was added methylisocyanate (1.1 mL, 18.0 mmol). After stirring at ambient temperature for 48 hours, the reaction was concentrated *in vacuo* and purified by silica gel chromatography (40% ethyl acetate in hexane) to afford the title compound (460 mg, 83%) as a white foam.

15 ¹H NMR (DMSO-*d*₆): δ 0.93 (3H, s), 0.96 (3H, s), 1.21 (9H, s), 1.58 (2H, t), 2.40-2.50 (1H, m), 2.54 (3H, d), 2.72-2.97 (3H, m), 3.24-3.42 (3H, m), 3.65-3.75 (1H, m), 3.82 (3H, s), 3.99 (2H, t), 4.99 (1H, d), 6.62 (1H, d), 6.88 (1H, q), 7.06 (2H, d), 7.10-7.24 (5H, m), 7.72 (2H, d) ; MS: 608 (MH⁺).

20

Step 6:



-191-

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-4-*N'*-methylcarbamoyloxybutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

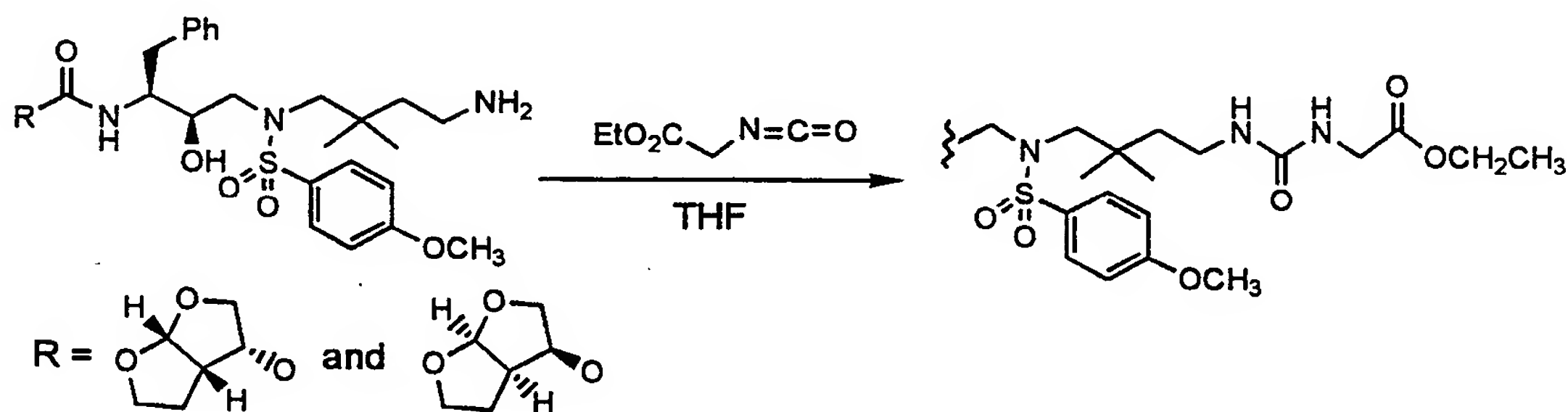
The product from step 5 was subjected to a protocol similar to that used in Example (Compound 210) to give the title compound as a foam.

¹H NMR (DMSO-*d*₆): δ 0.93 (1H, s), 0.94 (1H, s), 1.2 (1H, dd), 1.3-1.42 (1H, m), 1.6 (2H, t), 2.4 (1H, t), 2.55 (3H, d), 2.7-2.8 (2H, m), 2.83-3.00 (2H, m), 3.20-3.45 (3H, m), 3.55-3.62 (2H, m), 3.65-3.80 (3H, m), 3.82 (3H, s), 4.0 (2H, s), 4.8 (1H, quartet), 5.1 (1H, d), 5.5 (1H, d), 6.9 (1H, br s), 7.1 (2H, d), 7.15-7.25 (6H, m), 7.78 (2H, d); MS: 664 (MH⁺)

15

Example (Compound 222)

Step 1:



20

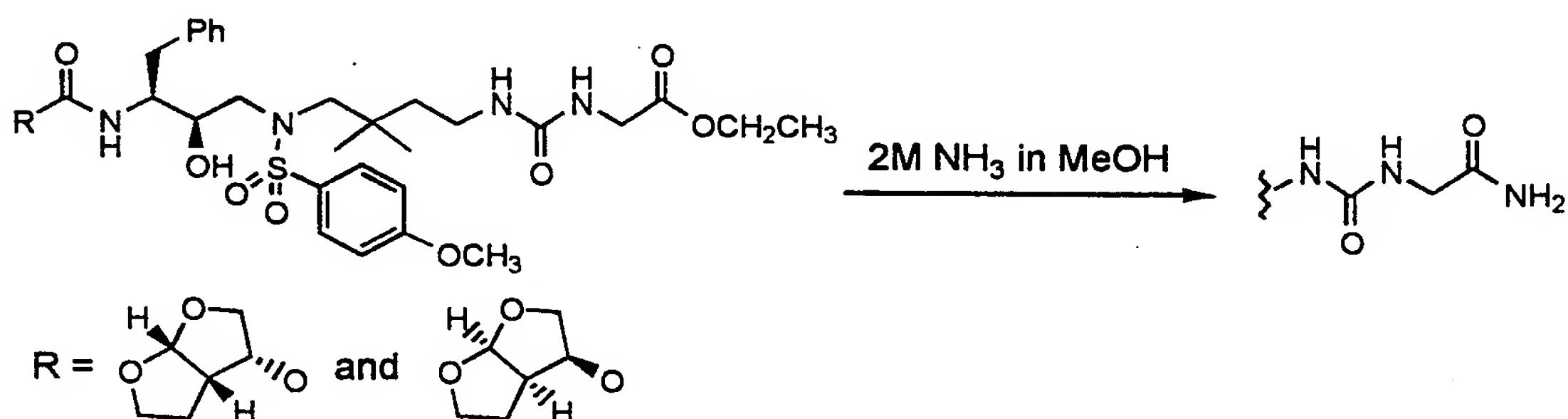
Ethyl (3*S*,4*R*)-1-[(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl)sulfonyl]-8,8-dimethyl-1,12-dioxo-2,6,11,13-tetraazapentadecan-15-oate and ethyl (3*S*,4*R*)-1-[(3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl)sulfonyl]-8,8-dimethyl-1,12-dioxo-2,6,11,13-tetraazapentadecan-15-oate

-192-

A solution of 50 mg (0.083 mmol) of a 1:1 mixture of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and
 5 (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate (Example (Compound 217)) in 3 mL of anhydrous THF was treated with 9.0 μ L (0.087 mmol) of ethyl isocyanato acetate and the
 10 resulting solution stirred at RT. After 2.5 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 47 mg (77%) of the desired product as a white foam.

15 ¹H NMR (CDCl₃): δ 7.70 (2H), 7.28-7.11 (5H), 7.00 (2H), 5.62 (1H), 5.47-5.30 (1H), 5.02 (1H), 4.24-3.46 (13H), 3.37-2.58 (9H), 2.04-1.39 (4H), 1.24 (3H), 1.03 (3H), 0.88 (3H). MS (ESI): 735 (M+H).

20 Step 2:



(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-[4-((2-amino-2-oxoethyl)amino)carbonylamino]-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and

25 (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-

-193-

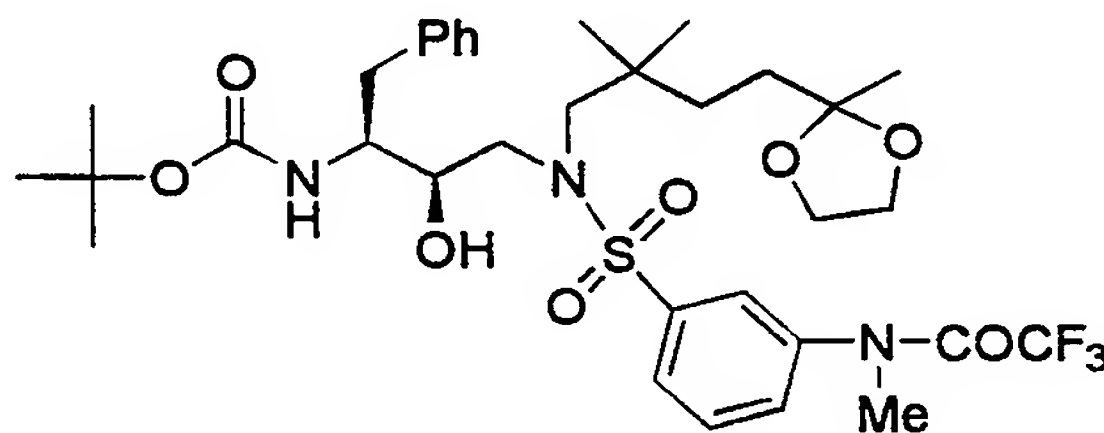
[4-([(2-amino-2-oxoethyl) amino] carbonylamino) -2,2-dimethylbutyl] [(4-methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate

A solution of 22 mg (0.030 mmol) of a 1:1 mixture of ethyl (3*S*,4*R*)-1-[(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl) sulfonyl]-8,8-dimethyl-1,12-dioxo-2,6,11,13-tetraazapentadecan-15-oate and ethyl (3*S*,4*R*)-1-[(3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl) sulfonyl]-8,8-dimethyl-1,12-dioxo-2,6,11,13-tetraazapentadecan-15-oate in 3 mL of 2M NH₃ in MeOH in a sealed tube was stirred at RT. After 5 days the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 93:7 CH₂Cl₂/2M NH₃ in MeOH) to afford 11 mg (52%) of the desired product as a white foam.

¹H NMR (CDCl₃): δ 7.69 (2H), 7.28-7.10 (5H), 6.97 (2H), 6.82-6.50 (1H), 6.12-4.43 (6H), 4.05 (1H), 3.97-3.42 (10H), 3.33-2.61 (9H), 2.00-1.35 (4H), 1.00 (3H), 0.88 (3H). MS (ESI): 706 (M+H).

Example (Compound 224)

Step 1:



ter-Butyl N-((1*S*,2*R*)-1-benzyl-3-[2,2-dimethyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl] [(3-(*N*-methyl,*N*-

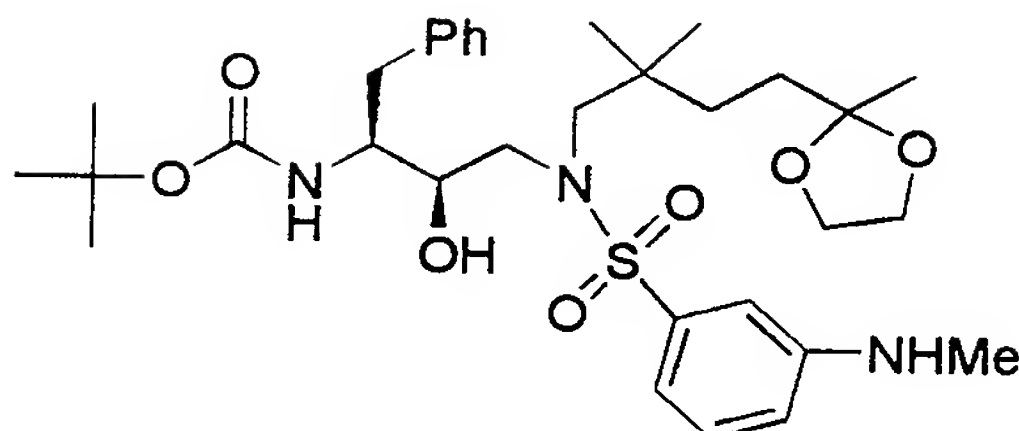
-194-

trifluoroacetyl)aminophenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

The product from Step 3 (Example (Compound 208)) (0.44 g, 0.8 mmol), was dissolved, in
5 dichloromethane (17 mL), and pyridine (1 mL, 0.97 g, 12.3 mmol) at 0°C was added trifluoroacetic anhydride (0.45 mL, 0.67 g, 3.2 mmol) and the mixture was stirred at ambient temperature for 2 hours. The mixture was washed with saturated sodium bicarbonate, dried (sodium sulfate) and
10 coevaporated with ethyl acetate (3X). The residue was dissolved in methanol (10 mL) and potassium carbonate (10 mg) was added. After stirring for 1 hour at ambient temperature, the solvent was evaporated and the residue was dissolved in acetone (15 mL) and potassium carbonate
15 (0.26 g, 2 mmol) and methyl iodide (0.25 mL, 0.56 g, 4.0 mmol) were added. After 3 hours, the solvent was evaporated and the residue was partitioned between dichloromethane and water. The organic phase was dried (sodium sulfate), evaporated, dried.

20 ¹H NMR (DMSO-d₆): δ 0.83 (6H, s), 1.2 (9H, s), 1.21-1.35 (2H, m), 1.5 (2H, br d), 2.40-2.48 (2H, m), 2.7-3.0 (3H, m), 3.05-3.15 (1H, m), 3.3-3.5 (8H, m), 3.65-3.75 (1H, m), 3.8 (3H, s), 4.9 (1H, d), 7.1-7.3 (6H, m), 7.6-7.8 (2H, m), 7.87 (1H, d), 7.95 (1H, s); MS: 738 (M + 23).

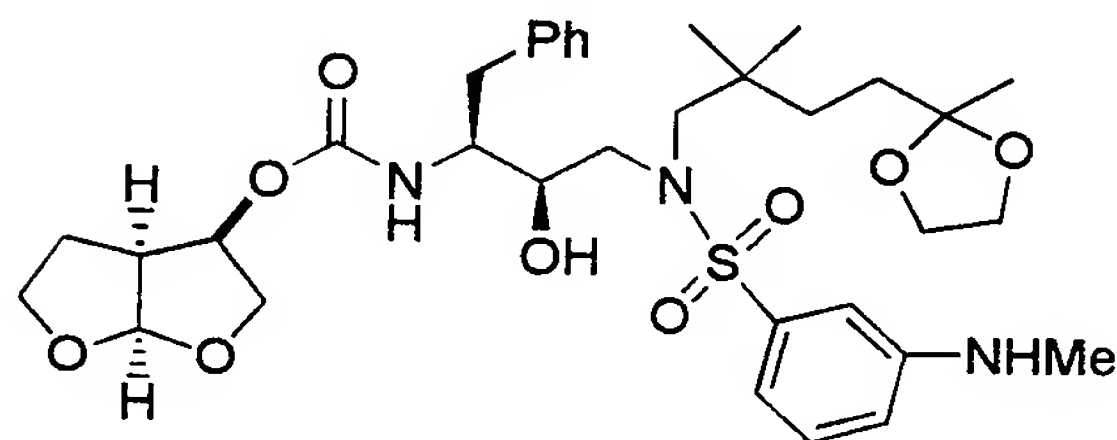
25 Step 2:



The product from Step 1 (0.16 g) was dissolved in 2M ammonia/ethanol (15 mL) and the mixture was stirred at ambient temperature for 2 hours. Solvent was evaporated and the residue was partitioned between water and dichloromethane. The organic phase was dried (sodium sulfate) and evaporated to give the title compound as a foam.

¹H NMR (CDCl₃): δ 0.92 (3H, s), 0.96 (3H, s), 1.2-1.4 (11, m), 1.6 (2H, d), 2.7-2.9 (6H, m), 3.0-3.3 (4H, m), 3.7 (1H, br s), 3.9 (7H, br s), 4.2 (1H, br s), 4.83 (1H, d), 6.75 (1H, d), 6.95 (1H, s), 7.02 (1H, d), 7.1-7.3 (7H, m); MS: 620 (MH⁺).

Step 3:

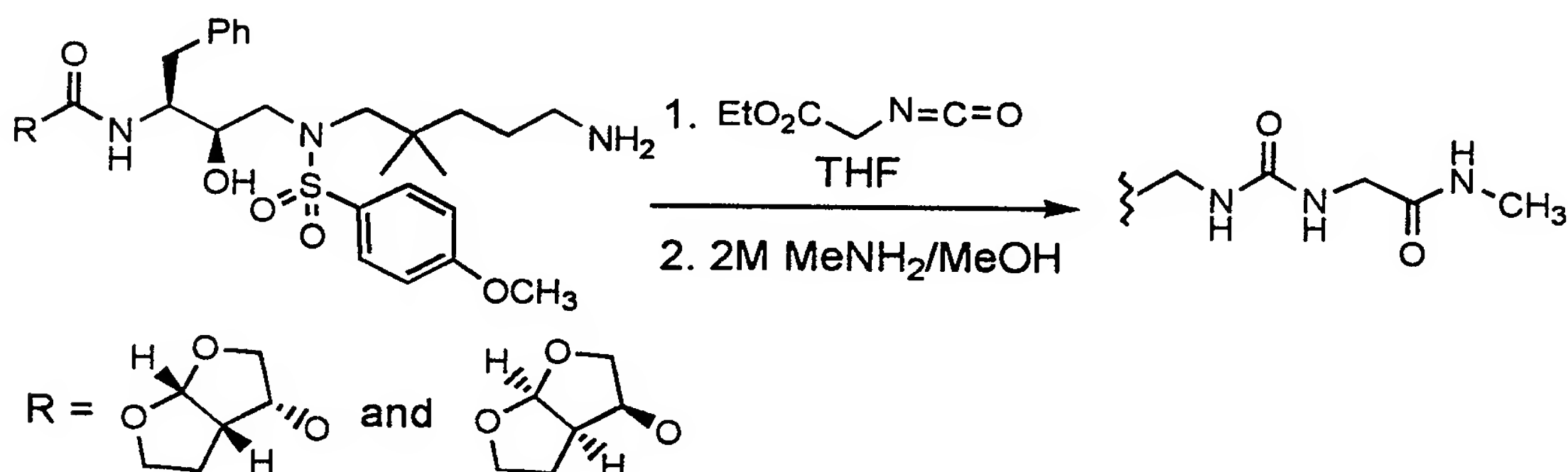


(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(2,2-dimethyl-5-oxohexyl)[(3-methylaminophenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

The product from Step 2 was subjected to the procedure used in Example (Compound 210) to afford the title compound as a foam.

¹H NMR (DMSO-d₆): δ 0.9 (3H, s), 0.94 (3H, s), 1.17 (1H, dd), 1.3-1.4 (1H, m), 1.45 (2H, t), 2.06 (3H, s), 2.35-2.45 (3H, m), 2.7 (3H, d), 2.71-2.80 (2H, m), 2.78 (1H, dd), 2.90 (1H, dd), 3.25-3.39 (3H, m), 3.4-3.5 (1H, m), 3.52-3.60 (2H, m), 3.7 (1H, t), 3.8 (2, dd), 4.8 (1H, quartet), 5.1 (1H, d), 5.5 (1H, d), 6.2 (1H, quartet), 6.75 (1H, d), 6.85 (1H, s), 6.86 (1H, d), 7.1-7.25 (6H, m); MS: 632 (MH⁺).

Example (Compound 225)

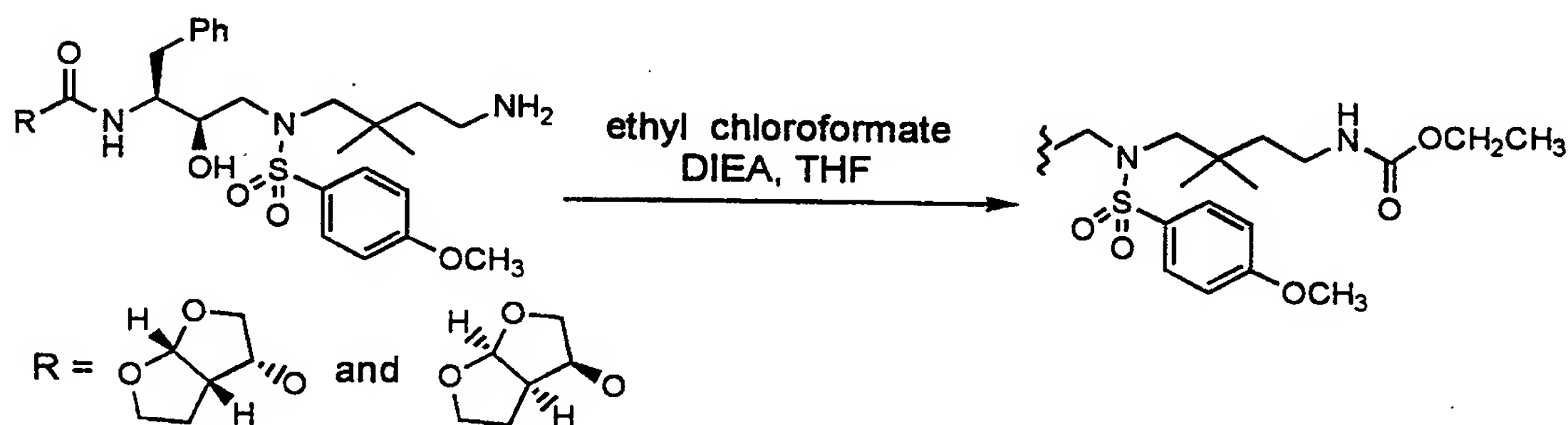


(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-2-hydroxy-4-[(4-methoxyphenyl)sulfonyl]-6,6-dimethyl-11,14-dioxo-4,10,12,15-tetraazahexadec-1-ylcarbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-2-hydroxy-4-[(4-methoxyphenyl)sulfonyl]-6,6-dimethyl-11,14-dioxo-4,10,12,15-tetraazahexadec-1-ylcarbamate

A solution of 39 mg (0.063 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)

-197-

- sulfonyl]amino-1-benzyl-2-hydroxypropyl) carbamate and (3*S*, 3*aR*, 6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-3-(5-amino-2,2-dimethylpentyl) [(4-methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate (Example
- 5 (Compound 212)) in 2 mL of anhydrous THF was treated with 7.4 μ L (0.066 mmol) of ethyl isocyanato acetate and the resulting solution was stirred at RT. After 2.5 hours the solution was concentrated *in vacuo*. The residue was dissolved in 3 mL of 2M MeNH₂ in MeOH in a sealed tube and
- 10 the solution was stirred at RT. After 18 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 93:7 CH₂Cl₂/2M NH₃ in MeOH) to afford 25 mg (54%) of the desired product as a white foam.
- 15 ¹H NMR (CDCl₃): δ 7.78 (2H), 7.33-7.15 (5H), 7.08-6.83 (3H), 6.30-4.80 (5H), 4.18-3.48 (11H), 3.38-2.62 (12H), 2.03-1.17 (6H), 0.98 (3H), 0.90 (3H). MS (ESI): 734 (M+H).

Example (Compound 226)

- (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-1-benzyl-3-4-[(ethoxycarbonyl) amino]-2,2-dimethylbutyl [(4-methoxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate
- 20 and (3*S*, 3*aR*, 6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-1-benzyl-3-4-[(ethoxycarbonyl) amino]-2,2-

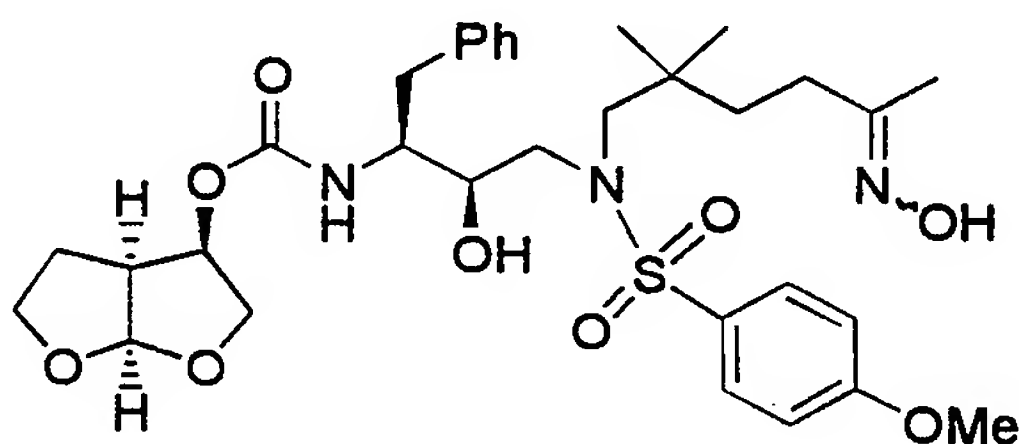
-198-

dimethylbutyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

A solution of 30 mg (0.050 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate (Example (Compound 217)) in 2 mL of anhydrous THF was cooled to 0°C and was treated with 9.6 µL (0.055 mmol) of N,N-diisopropylethylamine followed by 5.2 µL (0.055 mmol) of ethyl chloroformate. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated in vacuo and the residue was subjected to flash chromatography (SiO₂, 97:3 CH₂Cl₂/2M NH₃ in MeOH) to afford 31 mg (92%) of the desired product as a white foam.

¹H NMR (CDCl₃): 7.74 (2H), 7.35-7.16 (5H), 7.02 (2H), 5.68 (1H), 5.31 (1H), 5.18-4.90 (2H), 4.28-3.50 (11H), 3.35-2.70 (9H), 2.08-1.32 (4H), 1.27 (3H), 1.03 (3H), 0.98 (3H). MS (ESI): 678 (M+H).

Example (Compound 227)



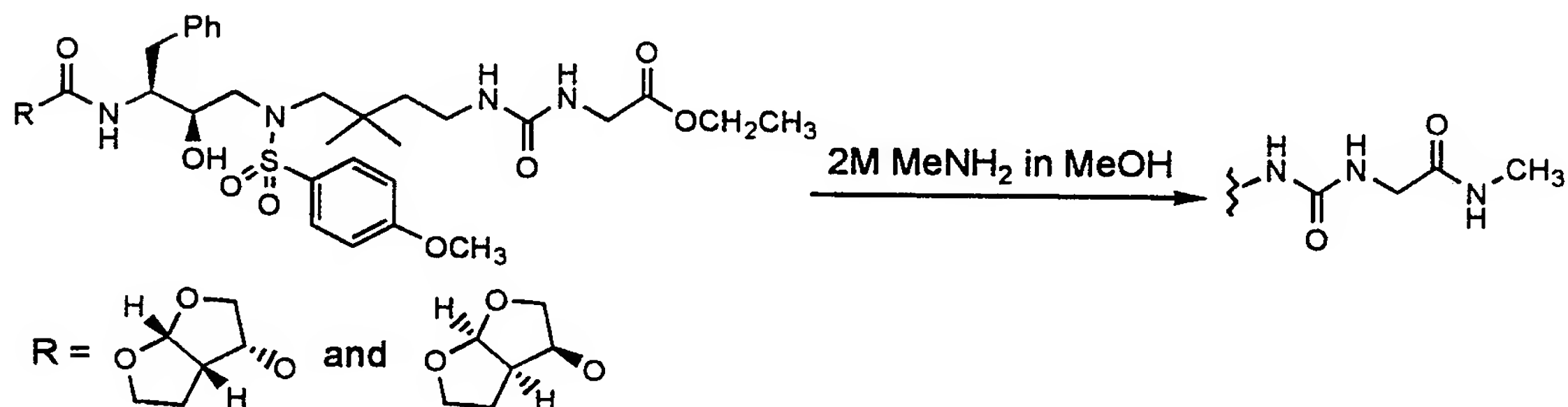
-199-

(3*R*,3*aS*,6*aR*) -Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-5-hydroxyiminohexyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

To a solution of the product from Example
 5 (Compound 210), (37 mg, 0.06 mmol) in pyridine (0.5 mL) and ethanol (0.5 mL) was added hydroxylamine hydrochloride (22 mg, 0.31 mmol) and the mixture was stirred at ambient temperature for 2 hours. Solvent was evaporated and the residue was dissolved in
 10 dichloromethane, washed with saturated sodium bicarbonate, dried (sodium sulfate), evaporated and chromatographed (silica gel, hexanes/ethyl acetate, 2:3) to provide the title compound as a foam; ¹H NMR (CDCl₃, signals for the major diastereomer): δ 0.85 (3H, s), 0.98 (3H, s), 1.3-1.7 (4H, m), 1.85 (3H, s), 2.1 (1H, dd),
 15 2.58 (1H, d), 2.7-3.0 (4H, m), 3.1-3.3 (3H, m), 3.6-3.7 (3H, m), 3.8-4.0 (3H, m), 3.9 (3H, s), 4.05-4.15 (2H, m), 5.0 (1H, quartet), 5.6 (2H, d), 7.0 (2H, d), 7.15-7.25 (5H, m), 7.75 (2H, d); MS: 648 (MH⁺); C₃₂H₄₅N₃O₉S.

20

Example (Compound 228)



(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-1-benzyl-2-hydroxy-4-[(4-methoxyphenyl)sulfonyl]-6,6-

-200-

dimethyl-10,13-dioxo-4,9,11,14-tetraazapentadec-1-ylcarbamate and (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-1-benzyl-2-hydroxy-4-[(4-methoxyphenyl)sulfonyl]-6,6-dimethyl-10,13-dioxo-4,9,11,14-tetraazapentadec-1-ylcarbamate

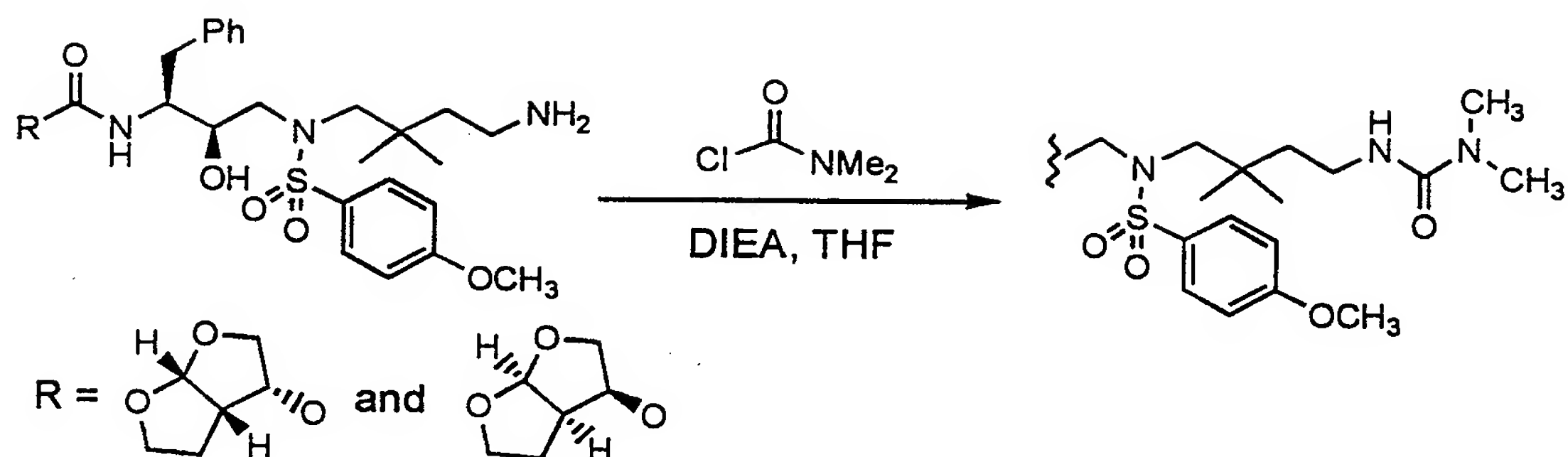
A solution of 22 mg (0.030 mmol) of a 1:1 mixture of ethyl (3*S*,4*R*)-1-[(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl)sulfonyl]-8,8-dimethyl-1,12-dioxo-2,6,11,13-tetraazapentadecan-15-oate and ethyl (3*S*,4*R*)-1-[(3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl)sulfonyl]-8,8-dimethyl-1,12-dioxo-2,6,11,13-tetraazapentadecan-15-oate in 3 mL of 2*M* methylamine in MeOH in a sealed tube was stirred at RT. After 18 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 93:7 CH₂Cl₂/2*M* NH₃ in MeOH) to afford 13 mg (60%) of the desired product as a white foam.

¹H NMR (CDCl₃): 7.75 (2H), 7.37-7.13 (5H), 7.02 (2H), 6.90-6.61 (1H), 6.10-4.49 (5H), 4.15-3.50 (11H), 3.40-2.67 (12H), 2.10-1.20 (4H), 1.06 (3H), 0.93 (3H).

MS (ESI): 720 (M+H).

Example (Compound 229)

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(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(4-[(dimethylamino)carbonyl]amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate and

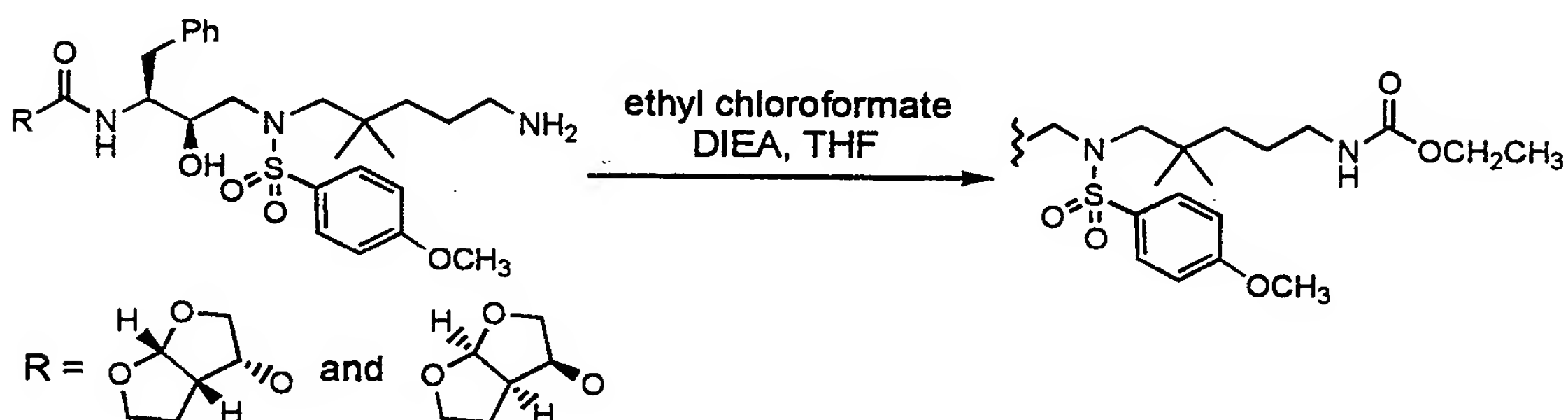
- 5 (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(4-[(dimethylamino)carbonyl]amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

A solution of 30 mg (0.050 mmol) of a 1:1
 10 mixture of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3*S*,3*aR*,6*aS*)hexahydrofuro
 [2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(4-amino-2,2-
 15 dimethylbutyl)[(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate (Example (Compound 217)) in 2 mL of anhydrous THF was cooled to 0°C and was treated with 9.6 µL (0.055 mmol) of *N,N*-diisopropylethylamine followed by 5.0 µL (0.053 mmol) of dimethylcarbamyl
 20 chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 30 mg (89%) of the desired product as a white foam.

-202-

^1H NMR (CDCl_3): 7.69 (2H), 7.29–7.10 (5H), 6.96 (2H), 5.80–5.52 (2H), 4.99–4.23 (2H), 3.99–3.43 (11H), 3.35–3.02 (5H), 3.01–2.62 (8H), 2.01–1.24 (4H), 1.01 (3H), 0.93 (3H). MS (ESI): 677 (M+H).

5

Example (Compound 230)

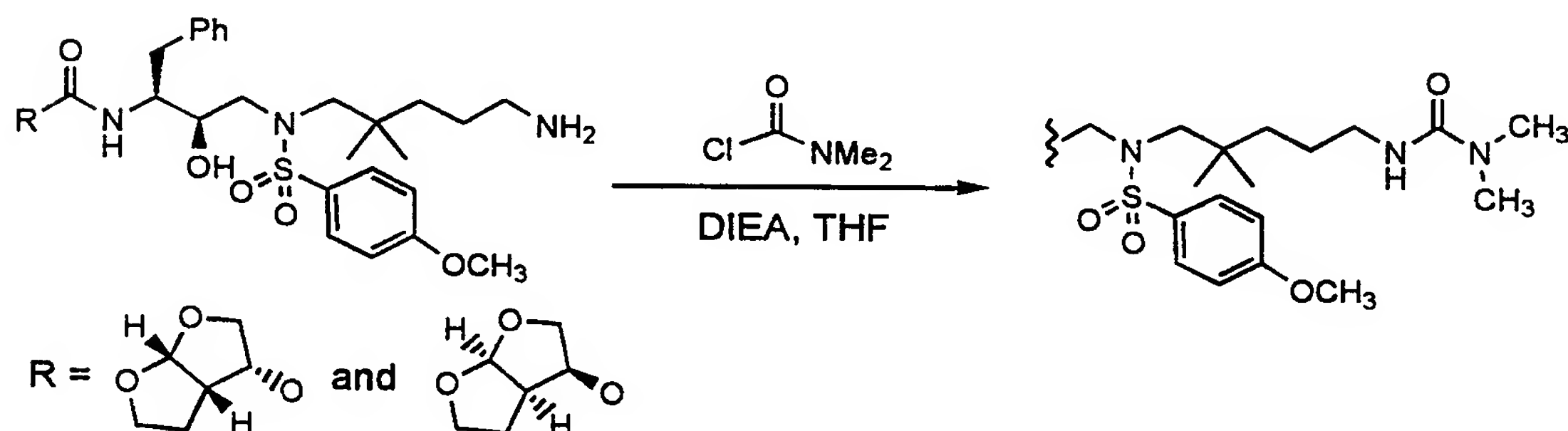
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-5-[(ethoxycarbonyl)amino]-2,2-dimethylpentyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-5-[(ethoxycarbonyl)amino]-2,2-dimethylpentyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

A solution of 30 mg (0.048 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl) [(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl) [(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate (Example (Compound 236)) in 2 mL of anhydrous THF at 0°C was treated with 9.2 μL (0.053 mmol) of N,N-diisopropylethylamine followed by 5.0 μL (0.053 mmol) of ethyl chloroformate. The resulting solution was allowed

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to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 97:3 CH₂Cl₂/2M NH₃ in MeOH) to afford 30 mg (94%) of the desired product as a white foam. H¹-NMR (CDCl₃): 7.76 (2H), 7.37-7.18 (5H), 7.04 (2H), 5.67 (1H), 5.60-5.32 (1H), 5.01 (2H), 4.24-3.44 (11H), 3.30-2.70 (9H), 2.10-1.20 (9H), 0.96 (3H), 0.90 (3H). MS (ESI): 692 (M+H).

Example (Compound 231)



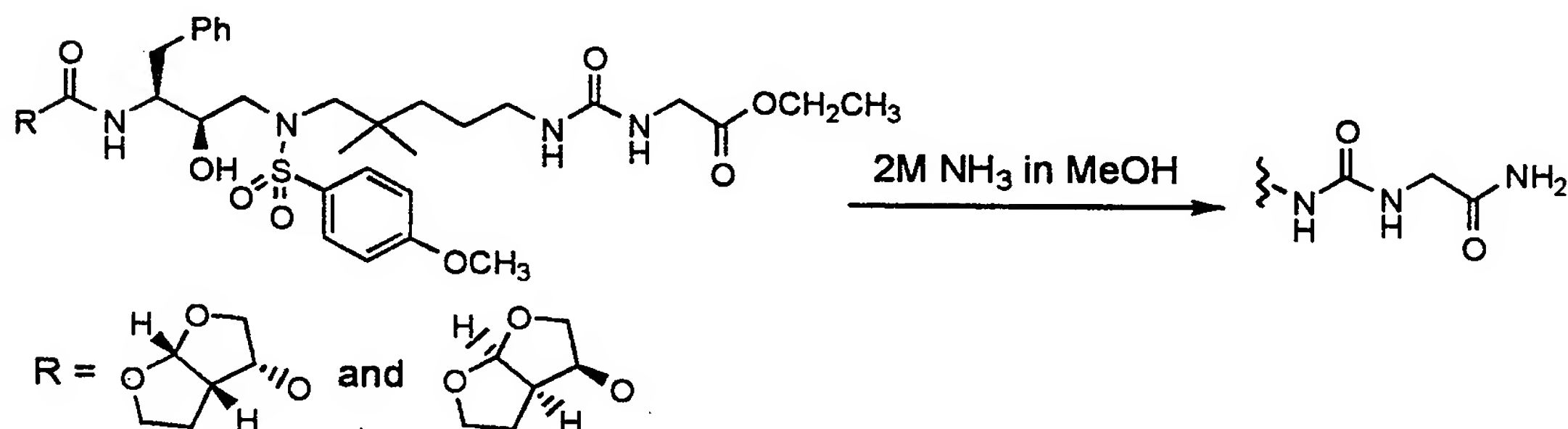
- (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(5-[(dimethylamino)carbonyl]amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate and
 (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(5-[(dimethylamino)carbonyl]amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

A solution of 30 mg (0.048 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl) carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-

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hydroxypropyl)carbamate (Example (Compound 212) in 2 mL of anhydrous THF at 0°C was treated with 9.2 μ L (0.053 mmol) of *N,N*-diisopropylethylamine followed by 4.6 μ L (0.050 mmol) of dimethylcarbamyl chloride. The resulting solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated in vacuo and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 26 mg (79%) of the desired product as a white foam. ¹H-NMR (CDCl₃): 7.71 (2H), 7.29-7.09 (5H), 6.98 (2H), 6.08-5.72 (1H), 5.61 (1H), 4.92 (1H), 4.65 (1H), 4.24 (1H), 3.99-3.43 (8H), 3.28-2.63 (15H), 1.99-1.70 (1H), 1.54-1.18 (5H), 0.92 (3H), 0.84 (3H). MS (ESI): 691 (M+H).

Example (Compound 232)



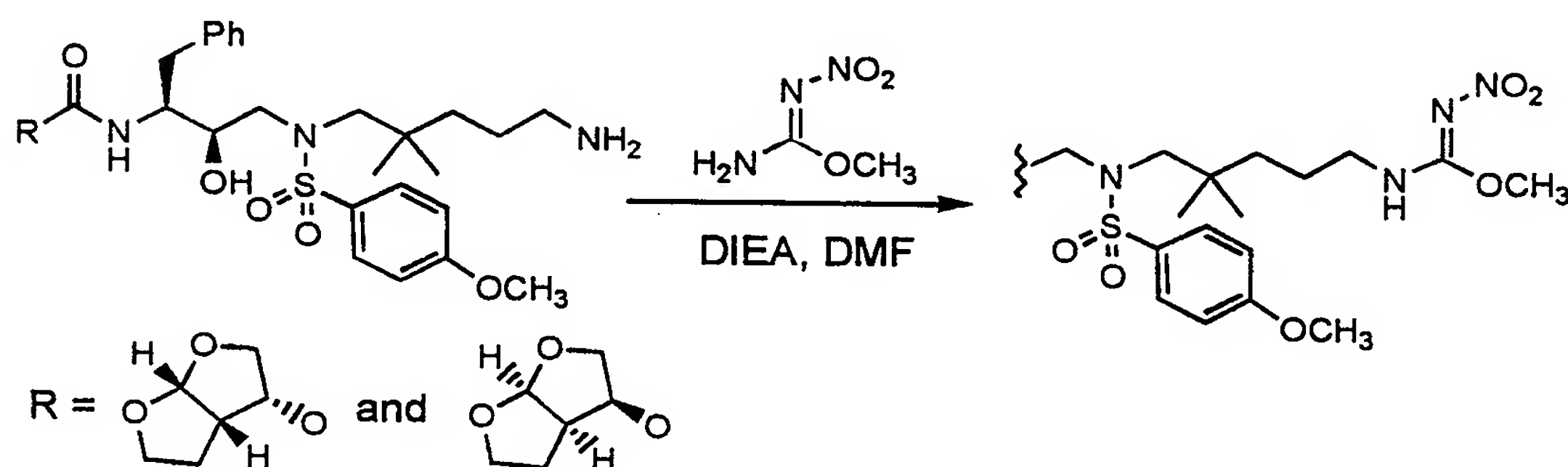
(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-[5-((2-amino-2-oxoethyl)amino)carbonylamino)-2,2-dimethylpentyl][(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-[5-((2-amino-2-oxoethyl)amino)carbonylamino)-2,2-dimethylpentyl][(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate

A solution of 27 mg (0.036 mmol) of a 1:1 mixture of ethyl (3*S*,4*R*)-1-

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[(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl) sulfonyl]-8,8-dimethyl-1,13-dioxo-2,6,12,14-tetraazahexadecan-16-oate and ethyl (3*S*, 4*R*)-1-[(3*S*, 3*aR*, 6*aS*) hexahydrofuro[2,3-*b*]furan-3-yloxy]-3-benzyl-4-hydroxy-6-[(4-methoxyphenyl) sulfonyl]-8,8-dimethyl-1,13-dioxo-2,6,12,14-tetraazahexadecan-16-oate (Example (Compound 216)) in 6 mL of 2M NH₃ in MeOH in a sealed tube was stirred at RT. After 3 days the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 93:7 CH₂Cl₂/2M NH₃ in MeOH) to afford 11 mg (43%) of the desired compound as a white foam. H¹-NMR (CDCl₃): 7.64 (2H), 7.29-7.01 (5H), 6.99-6.63 (3H), 6.40-5.80 (2H), 5.74-5.47 (2H), 5.00-4.74 (1H), 4.28-3.40 (12H), 3.28-2.42 (9H), 1.97-1.08 (6H), 0.89 (3H), 0.79 (3H). MS (ESI): 720 (M+H)

Example (Compound 233)



2-[(*Z*)-[(5-[(2*R*, 3*S*)-3-[(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yloxy]carbonylamino)-2-hydroxy-4-phenylbutyl][(4-methoxyphenyl) sulfonyl]amino-4,4-dimethylpentyl]amino] (methoxy)methylidene]-1-hydroxy-1-oxohydrazinium and 2-[(*Z*)-[(5-[(2*R*, 3*S*)-3-[(3*S*, 3*aR*, 6*aS*) hexahydrofuro[2,3-*b*]furan-3-

-206-

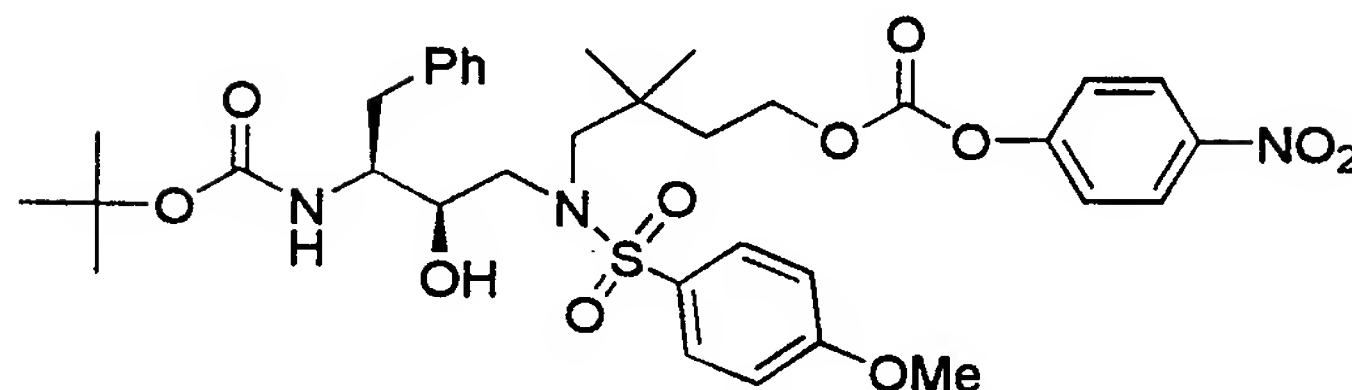
ylloxy]carbonylamino)-2-hydroxy-4-phenylbutyl][(4-methoxyphenyl)sulfonyl]amino-4,4-dimethylpentyl)amino] (methoxy)methylidene]-1-hydroxy-1-oxohydrazinium

- 5 A solution of 30 mg (0.048 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate
- 10 and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate (Example (Compound 212)) in 1.5 mL of anhydrous DMF was treated with 10 μ L (0.059 mmol) of N,N-diisopropylethylamine
- 15 followed by 7.0 mg (0.059 mmol) of O-methyl-N-nitroisourea and the resulting solution was stirred at RT. After 20 hours the solution was concentrated in vacuo and the residue subjected to flash chromatography (SiO₂, 93:7 CH₂Cl₂/2M NH₃ in MeOH) to afford 19 mg (56%) of
- 20 a white foam. ¹H-NMR (CDCl₃): 7.75 (2H), 7.37-7.13 (5H), 7.02 (2H), 5.75-5.23 (2H), 5.01 (1H), 4.20-3.48 (12H), 3.25-2.62 (9H), 2.10-1.75 (1H), 1.65-1.30 (5H), 0.98 (3H), 0.93 (3H).

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Example (Compound 234)

Step 1:

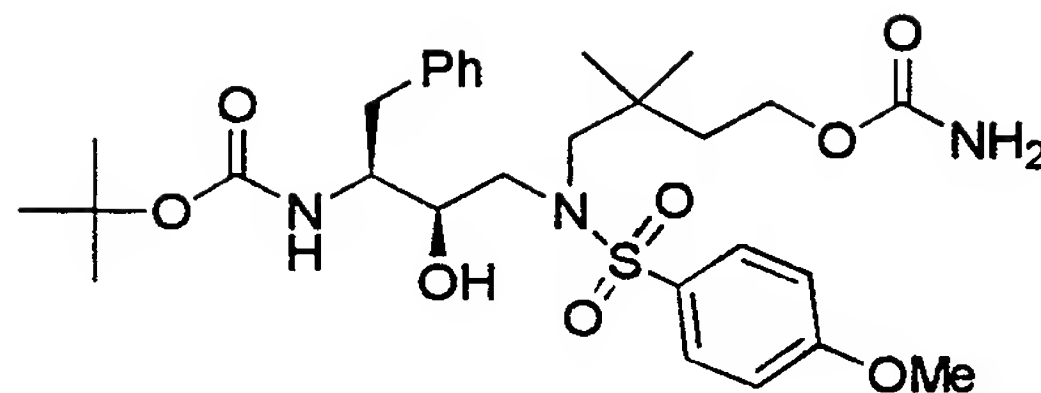


ter-Butyl N-((1S,2R)-1-benzyl-3-(2,2-dimethyl-4-[(4-nitrophenoxy)carbonyloxy]butyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

To a solution of *p*-nitrophenyl chloroformate (320 mg, 1.6 mmol) in dichloromethane (8 mL) at 0°C was added pyridine (2 mL) followed by a solution of the product from step 4, Example (Compound 221), (500 mg, 0.91 mmol) in dichloromethane (7 mL). The mixture was stirred at ambient temperature for 18 hours. The mixture was diluted with ethyl ether and washed with 0.5 N hydrochloric acid, saturated sodium bicarbonate, brine, dried (sodium sulfate), and evaporated. The residue was chromatographed (silica gel, hexanes/ethyl acetate, 7:3) to afford the title compound as a white foam (530 mg); ¹H NMR (DMSO-*d*₆): δ 0.97 (3H, s), 0.99 (3H, s), 1.2 (9H, s), 1.89 (2H, t), 2.40-2.46 (1H, m), 2.78 (1H, d), 2.88-3.00 (2H, m), 3.3-3.4 (3H, m), 3.75 (1H, br quartet), 3.8 (3H, s), 4.38 (2H, t), 5.05 (1H, d), 6.65 (1H, d), 7.06 (2H, d), 7.10-7.22 (5H, m), 7.55 (2H, d), 7.75 (2H, d), 8.3 (2H, d)

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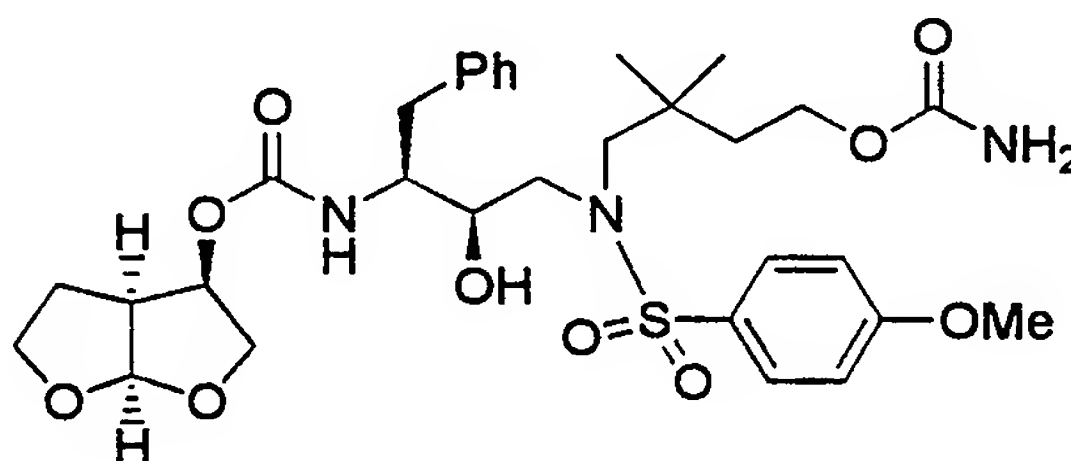
Step 2:



tert-Butyl N-((1S,2R)-1-benzyl-3-(4-carbamoyloxy-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

- 5 To a solution of the product from step 1 (520 mg, 0.73 mmol) in tetrahydrofuran (10 mL) at 5°C was added ammonium hydroxide (30%, 2 mL) and the reaction was stirred at 5°C for 40 minutes. The mixture was diluted with ethyl ether, washed with saturated sodium
- 10 bicarbonate, dried (magnesium sulfate), and evaporated to provide the title compound (490 mg) containing p-nitrophenol (~10%). ¹H NMR (DMSO-d₆): δ 0.93 (3H, s), 0.96 (3H, s), 1.2 (9H, s), 1.6 (2H, t), 2.45 (1H, dd), 2.78 (1H, d), 2.85-2.95 (2H, m), 3.3-3.6 (3H, m), 3.7
- 15 (1H, quartet), 3.8 (3H, s), 3.96 (2H, t), 5.0 (1H, d), 6.4 (2H, br s), 6.6 (1H, d), 7.05 (2H, d), 7.10-7.25 (5H, m), 7.7 (2H, d).

Example (Compound 235)

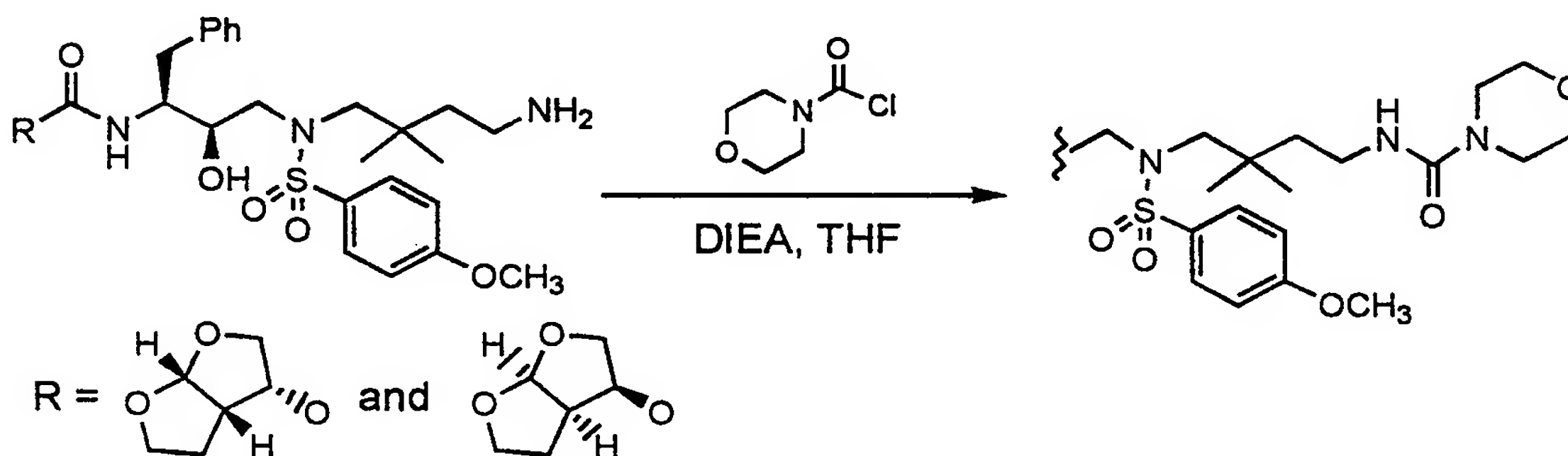


- 20 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(4-carbamoyloxy-2,2-dimethylbutyl)[(4-

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methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

The product from Example (Compound 234) was subjected to a procedure described in Example (Compound 216) to afford the title compound as a foam; ¹H NMR (DMSO-
 5 d₆): δ 0.92 (3H, s), 0.96 (3H, s), 1.2 (1H, dd), 1.3-1.4 (1H, m), 1.6 (2H, t), 2.4 (1H, dd), 2.7-2.8 (2H, m), 2.9-3.0 (2H, m), 3.3-3.5 (3H, m), 3.55-3.65 (2H, m), 3.66-3.80 (2H, m), 3.8 (3H, s), 3.96 (2H, t), 4.8 (1H, quartet), 5.1 (1H, d), 5.5 (1H, d), 6.4 (2H, br s), 7.05
 10 (2H, d), 7.1-7.3 (6H, m), 7.7 (2H, d); MS: 672 (M + 23).

Example (Compound 236)

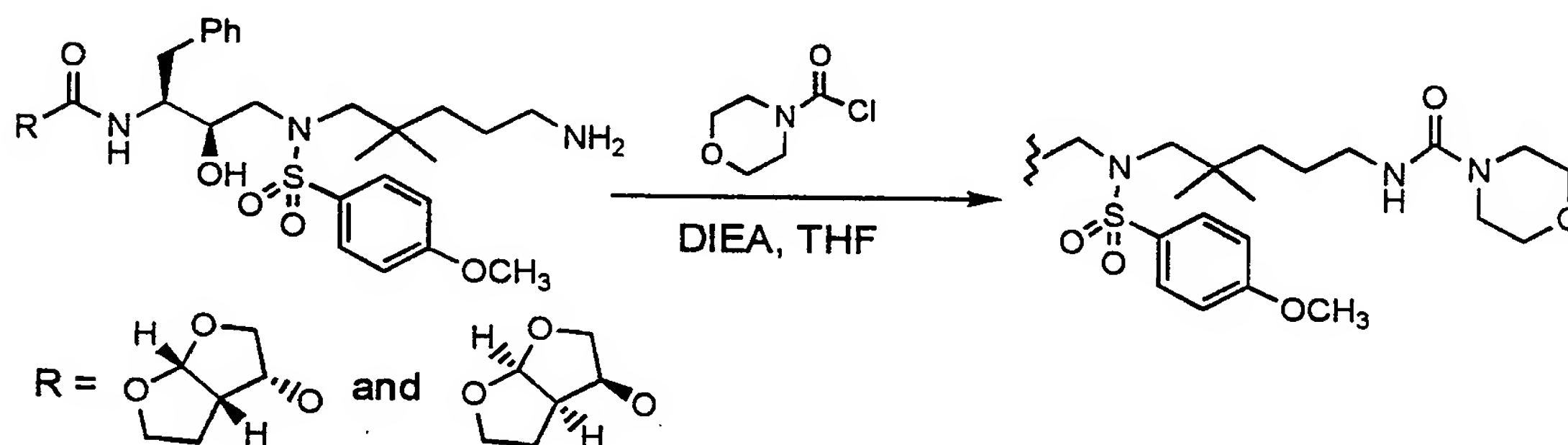
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-2,2-dimethyl-4-[(morpholinocarbonyl)amino]butyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate and (3S,3aR,6aS)hexahydrofuro
 15 [2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-2,2-dimethyl-4-[(morpholinocarbonyl)amino]butyl[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

A solution of 30 mg (0.050 mmol) of a 1:1
 20 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-

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(4-amino-2,2-dimethylbutyl) [(4-methoxyphenyl) sulfonyl] amino-1-benzyl-2-hydroxypropyl) carbamate in 1.5 mL of anhydrous THF was cooled to 0°C and was treated with 9.6 μL (0.055 mmol) of *N,N*-diisopropylethylamine followed by 6.4 μL (0.055 mmol) of 4-morpholinecarbonyl chloride. The solution was allowed to warm to RT with stirring. After 2 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 31 mg (86%) of the desired product as a white foam. ¹H-NMR (CDCl₃): 7.74 (2H), 7.37-7.13 (6H), 7.01 (2H), 5.68 (1H), 5.42 (1H), 5.00 (2H), 4.32 (1H), 4.11-3.50 (14H), 3.49-2.70 (11H), 2.10-1.22 (4H), 1.13-0.94 (6H). MS (ESI): 741 (M+Na).

Example (Compound 237)

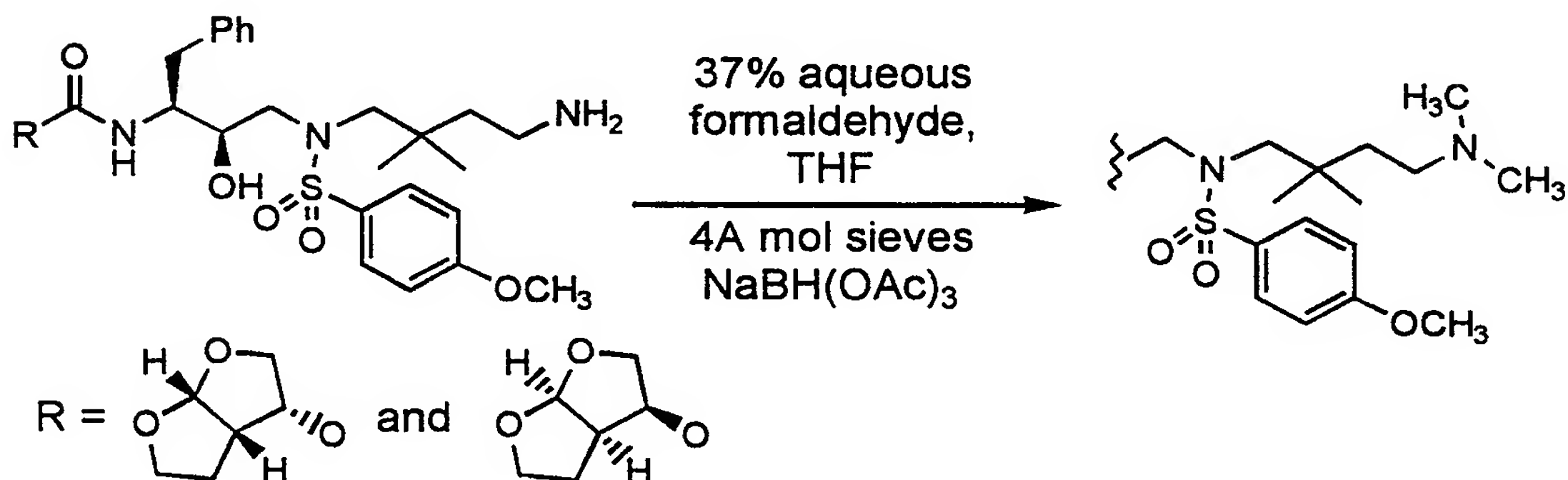


(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-2,2-dimethyl-5-[(morpholinocarbonyl) amino]pentyl [(4-methoxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate and 3*S*,3*aR*,6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-2,2-dimethyl-5-[(morpholinocarbonyl) amino]pentyl [(4-methoxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate

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A solution of 30 mg (0.048 mmol) of a 1:1 mixture of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate and (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate in 1.5 mL of anhydrous THF at 0°C was treated with 9.2 µL (0.053 mmol) of *N,N*-diisopropylethylamine followed by 6.2 µL (0.053 mmol) of 4-morpholinecarbonyl chloride. The resulting solution was allowed to warm to RT with stirring. After 2 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 31 mg (89%) of the desired product as a white foam. ¹H-NMR (CDCl₃): 7.74 (2H), 7.36-7.19 (6H), 7.02 (2H), 5.74-5.38 (2H), 4.98 (2H), 4.20 (1H), 4.12-3.46 (12H), 3.41-2.67 (13H), 2.04-1.22 (6H), 1.01-0.86 (6H). MS(ESI): 755 (M+Na).

20

Example (Compound 238)

(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-[4-(dimethylamino)-2,2-dimethylbutyl][(4-

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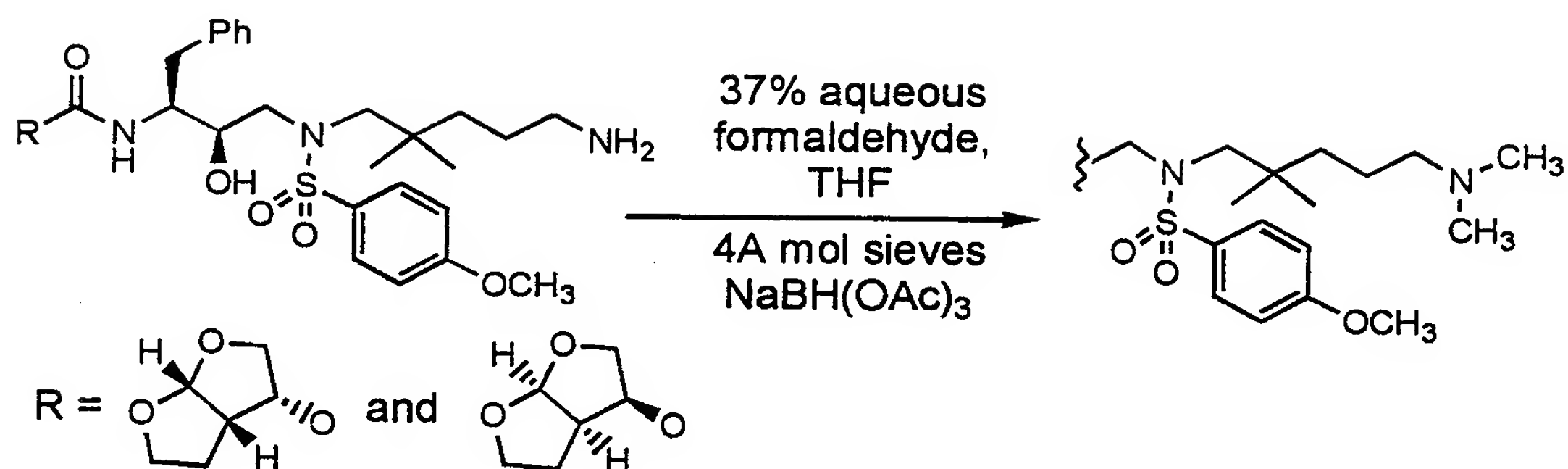
methoxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate
and (3*S*,3*aR*,6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-
((1*S*,2*R*)-1-benzyl-3-[4-(dimethylamino)-2,2-
dimethylbutyl] [(4-methoxyphenyl) sulfonyl] amino-2-
5 hydroxypropyl) carbamate

A solution of 50 mg (0.083 mmol) of a 1:1
mixture of (3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-
((1*S*,2*R*)-3-(4-amino-2,2-dimethylbutyl) [(4-
methoxyphenyl) sulfonyl] amino-1-benzyl-2-
10 hydroxypropyl) carbamate and
(3*S*,3*aR*,6*aS*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-
(4-amino-2,2-dimethylbutyl) [(4-methoxyphenyl) sulfonyl]
amino-1-benzyl-2-hydroxypropyl) carbamate and 0.034 mL
(0.42 mmol) of 37% aqueous formaldehyde in 5 mL of THF
15 was treated with 88 mg (0.42 mmol) of NaBH(OAc)₃ followed
by 150 mg of powdered 4A molecular sieves. The resulting
mixture was stirred at RT for 18 hours, filtered to
remove solids, and the filtrate concentrated in vacuo.
The residue was purified by flash chromatography (SiO₂,
20 95:5 to 9:1 CH₂Cl₂/2M NH₃ in MeOH) to afford 40 mg (75%)
of the desired compound as a white foam. . H¹-NMR (CDCl₃):
7.70 (2H), 7.30-7.11 (6H), 6.98 (2H), 5.62 (1H), 5.42-
5.22 (1H), 4.96 (1H), 4.09-3.44 (9H), 3.15-2.60 (7H),
2.38-1.30 (12H), 0.91 (6H). MS(ESI): 634 (M+H).

25

Example (Compound 239)

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(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-[5-(dimethylamino)-2,2-dimethylpentyl][(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate and (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-[5-(dimethylamino)-2,2-dimethylpentyl][(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

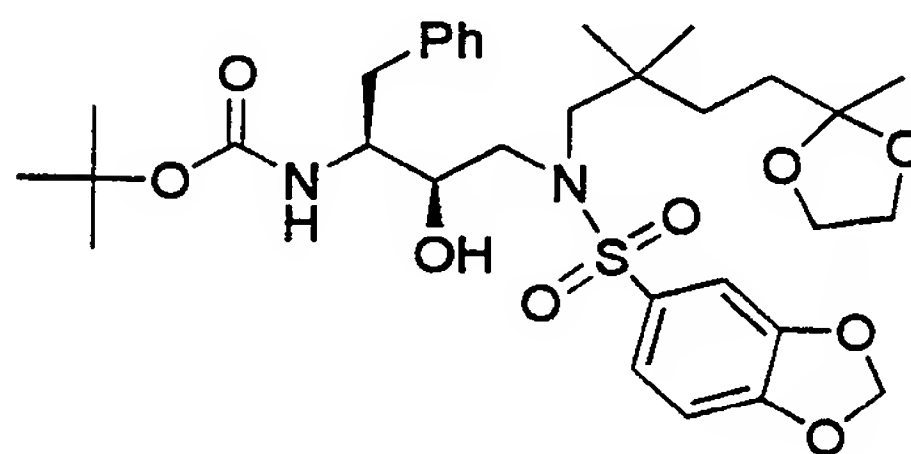
A solution of 50 mg (0.081 mmol) of a 1:1 mixture of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl][(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate and (3*S*,3*aR*,6*aS*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-3-(5-amino-2,2-dimethylpentyl][(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate and 0.033 mL (0.41 mmol) of 37% aqueous formaldehyde in 5 mL of THF was treated with 87 mg (0.41 mmol) of NaBH(OAc)₃ followed by 150 mg of powdered 4A molecular sieves. The resulting mixture was stirred at RT for 18 hours, filtered to remove solids, and the filtrate concentrated *in vacuo*. The residue was purified by flash chromatography (SiO₂, 9:1 CH₂Cl₂/2M NH₃ in MeOH) to afford 38 mg (73%) of the desired compound as a white foam. . H1-NMR (CDCl₃): 7.72 (2H), 7.30-7.10 (6H), 7.00 (2H), 6.98-6.40 (1H), 5.60 (1H), 4.92 (1H),

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4.00-3.07 (11H), 3.04-2.27 (5H), 2.18 (6H), 2.00-1.11 (8H), 1.03-0.81 (6H). MS (ESI): 648 (M+H).

Example (Compound 240)

Step 1:

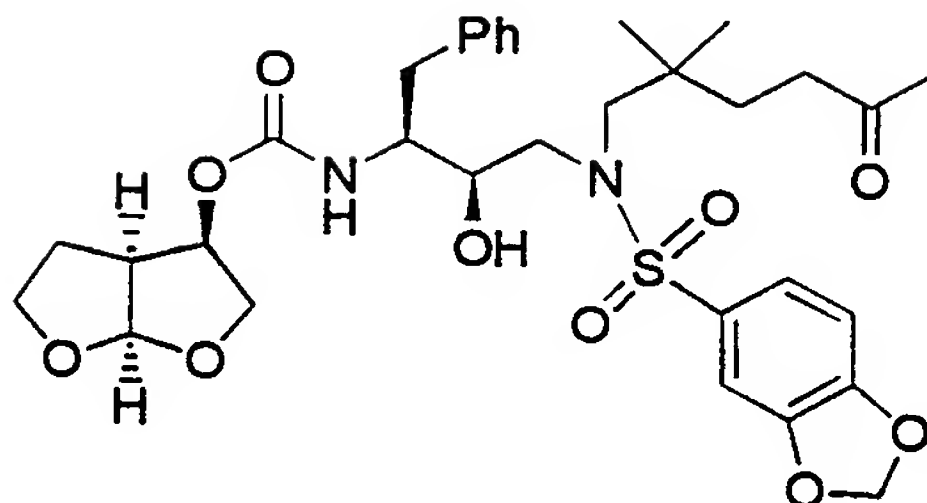


- 5 *tert*-Butyl N-((1*S*,2*R*)-1-benzyl-3-[2,2-dimethyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl][(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

The product from step 2 (Example (Compound
10 208)) was reacted with 3,4-methylenedioxyphenylsulfonyl chloride similar to procedure 3 (Example (Compound 208)) to provide the title compound as a foam; ¹H NMR (DMSO-*d*₆):
δ 0.85 (3H, s), 1.1-1.2 (1H, m), 1.22 (9H, s), 1.3 (2H, br dd), 1.5 (2H, br d), 2.2-2.4 (1H, m), 2.84 (1H, d),
15 2.9 (1H, dd), 2.97 (1H, dd), 3.30-3.45 (5H, m), 3.75 (1H, br quartet), 3.85 (4H, s), 5.0 (1H, d), 6.18 (2H, s), 6.6 (1H, d), 7.0 (1H, d), 7.1-7.3 (6H, m), 7.35 (1H, d); MS: 635 (MH⁺); C₃₂H₄₆N₂O₉S.

Step 2:

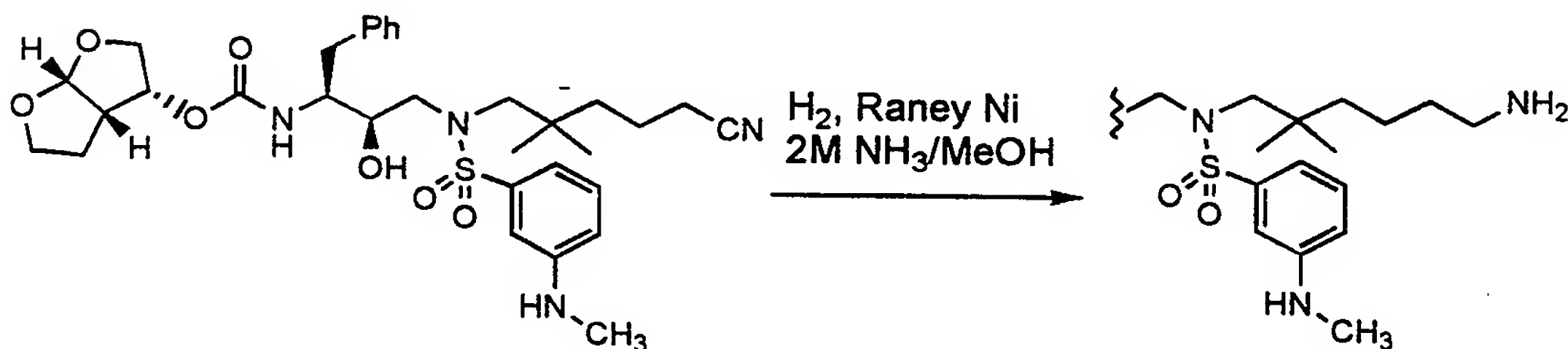
-215-



(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(2,2-dimethyl-5-oxohexyl)[(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate

5 The product from step a was subjected to procedure similar to Example (Compound 205) to afford the title compound as a foam; ^1H NMR (DMSO- d_6): δ 0.85 (3H, s), 0.90 (3H, s), 1.05 (1H, br dd), 1.2-1.4 (1H, m), 1.45 (2H, t), 2.1 (3H, s), 2.30-2.45 (3H, m), 2.73 (2H, br d), 2.9 (1H, dd), 2.95 (1H, d), 3.30-3.45 (3H, m), 3.5-3.6 (2H, m), 3.65-3.85 (3H, m), 4.8 (1H, quartet), 5.15 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 7.03 (1H, d), 7.1-7.25 (7H, m), 7.35 (1H, d); MS: 669 (M + 23); $\text{C}_{32}\text{H}_{42}\text{N}_2\text{O}_{10}\text{S}$.

Example (Compound 241)



15 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-((6-amino-2,2-dimethylhexyl)[3-(methylamino)phenyl]sulfonylamino)-1-benzyl-2-hydroxypropyl]carbamate

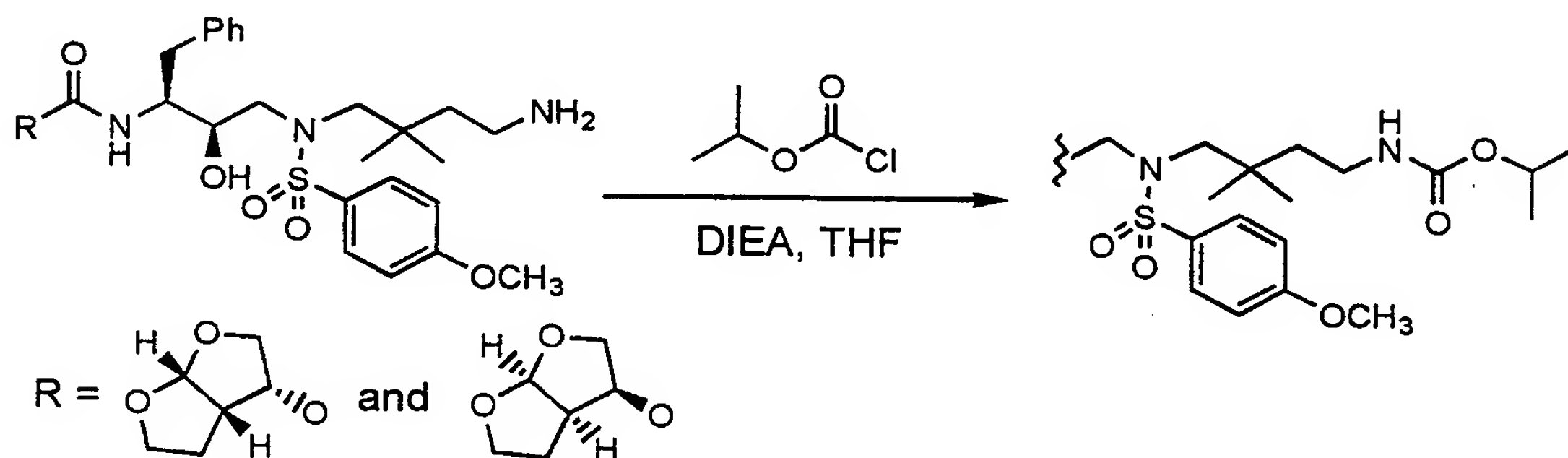
-216-

A mixture of 0.18 g (0.29 mmol) of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-benzyl-3-((5-cyano-2,2-dimethylpentyl)[3-(methylamino)phenyl]sulfonylamino)-2-hydroxypropyl]carbamate (obtained via a protocol directly analogous to Example (Compound 245)) and approximately 50 mg of Raney nickel (washed with water and then MeOH) in 40 mL of 2M NH₃/MeOH was subjected to hydrogenation at 40 psi. After 1.5 hours the vessel was purged with nitrogen, catalyst removed by filtration through celite and the filtrate concentrated to dryness in vacuo. The residue was subjected to flash chromatography (SiO₂, 9:1 CH₂Cl₂, 2M NH₃ in MeOH) to afford 0.13 g (72%) of the desired amine as a white foam.

¹H-NMR (CDCl₃): 7.31-7.12 (m, 6H), 7.01 (d, 1H), 6.92 (s, 1H), 6.75 (dd, 1H), 5.60 (d, 1H), 5.29 (d, 1H), 4.99 (q, 1H), 4.15 (br s, 1H), 4.00 (t, 1H), 3.89 (m, 1H), 3.80 (m, 2H), 3.67 (m, 2H), 3.21 (dd, 1H), 3.17-2.98 (m, 3H), 2.94-2.80 (m, 2H), 2.84 (s, 3H), 2.71 (m, 3H), 1.58 (m, 1H), 1.47-1.21 (m, 7H), 0.90 (s, 3H), 0.88 (s, 3H).

MS (ESI): 633 (M+H).

Example (Compound 242)



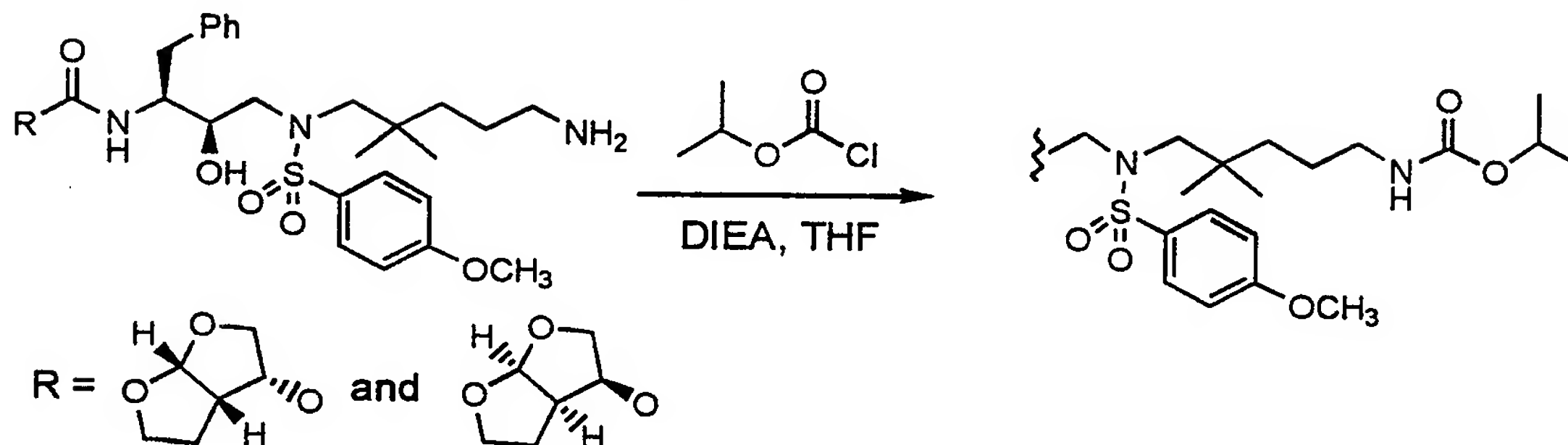
-217-

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-2-hydroxy-3-4-[(isopropoxycarbonyl)amino]-2,2-dimethylbutyl[(4-methoxyphenyl)sulfonyl]aminopropyl)carbamate and

- 5 (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-2-hydroxy-3-4-[(isopropoxycarbonyl)amino]-2,2-dimethylbutyl[(4-methoxyphenyl)sulfonyl]aminopropyl)carbamate.

- A solution of 71 mg (0.12 mmol) of a 1:1
10 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate and (3S,3aR,6aS)hexahydrofuro
[2,3-b]furan-3-yl N-((1S,2R)-3-(4-amino-2,2-
15 dimethylbutyl)[(4-methoxyphenyl)sulfonyl]amino-1-benzyl-2-hydroxypropyl)carbamate in 5 mL of anhydrous THF at 0°C was treated with 0.022 mL (0.13mmol) of N,N-diisopropylethylamine followed by 0.13 mL (0.13 mmol) of 1M isopropyl chloroformate/toluene solution. The
20 solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated in vacuo and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH) to afford 48 mg (58%) of the desired compound as a white foam. . H1-NMR (CDCl₃): 7.70 (2H), 7.30-7.14
25 (5H), 6.99 (2H), 5.62 (1H), 5.21 (1H), 5.01-4.81 (3H), 4.19-3.44 (10H), 3.27-2.68 (9H), 2.01-1.30 (4H), 1.21 (6H), 1.01-0.92 (6H). MS(ESI): 714(M+Na).

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Example (Compound 243)

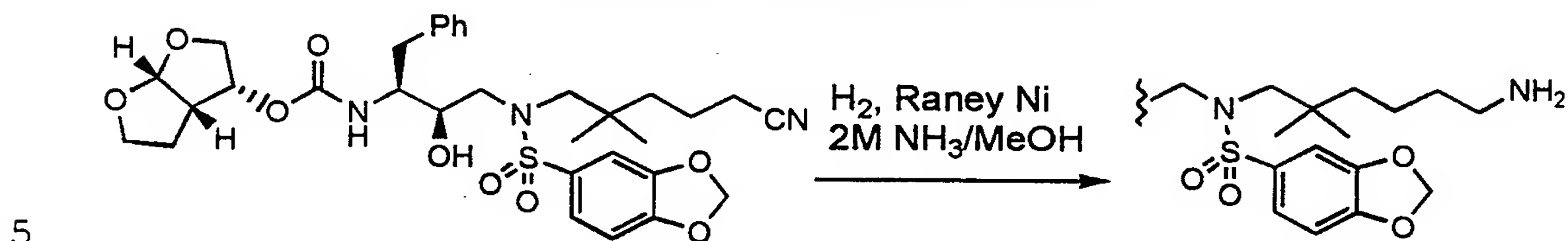
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-2-hydroxy-3-5-[(isopropoxycarbonyl)amino]-2,2-dimethylpentyl[(4-methoxyphenyl)sulfonyl]aminopropyl)carbamate and (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-2-hydroxy-3-5-[(isopropoxycarbonyl)amino]-2,2-dimethylpentyl[(4-methoxyphenyl)sulfonyl]aminopropyl)carbamate.

- 10 A solution of 70 mg (0.11 mmol) of a 1:1 mixture of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate and
- 15 (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(5-amino-2,2-dimethylpentyl)[(4-methoxyphenyl)sulfonyl] amino-1-benzyl-2-hydroxypropyl)carbamate in 5 mL of anhydrous THF at 0°C was treated with 0.022 mL (0.13mmol) of N,N-diisopropylethylamine followed by 0.12 mL (0.12
- 20 mmol) of 1M isopropyl chloroformate/toluene solution. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH) to afford 75 mg (95%) of the desired compound
- 25 as a white foam. . H1-NMR (CDCl₃): 7.70 (2H), 7.30-7.11

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(5H), 7.00 (2H), 5.65-5.26 (2H), 5.00-4.80 (3H), 4.18-3.42 (10H), 3.22-2.63 (9H), 2.00-1.18 (12H), 0.94-0.83 (6H). MS (ESI): 728 (M+Na).

Example (Compound 244)



(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-3-[(6-amino-2,2-dimethylhexyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate.

10 A mixture of 0.31 g (0.47 mmol) of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(5-cyano-2,2-dimethylpentyl)amino]-1-benzyl-2-hydroxypropylcarbamate (obtained via a protocol directly analogous to Example 15 (Compound 245)) and approximately 50 mg of Raney nickel (washed with water and then MeOH) in 40 mL of 2M NH₃/MeOH was subjected to hydrogenation at 45 psi. After 2.5 hours the vessel was purged with nitrogen, catalyst removed by filtration through celite and the filtrate 20 concentrated to dryness in vacuo. The residue was subjected to flash chromatography (SiO₂, 9:1 CH₂Cl₂, 2M NH₃ in MeOH) to afford 0.27 g (90%) of the desired amine as a white foam. H¹-NMR (CDCl₃): 7.33 (d, 1H), 7.28-7.13 (m, 6H), 6.87 (d, 1H), 6.08 (s, 2H), 5.60 (d, 1H), 5.31 (m, 1H), 5.00 (q, 1H), 4.00 (t, 1H), 3.91 (dd, 1H), 3.81 (m, 2H), 3.63 (m, 2H), 3.21-2.97 (m, 4H), 2.86 (m, 2H), 2.80-2.63 (m, 3H), 1.58 (m, 1H), 1.45-1.21 (m, 7H), 0.91 (d, 6H). MS (ESI): 648 (M+H).

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Example (Compound 245)

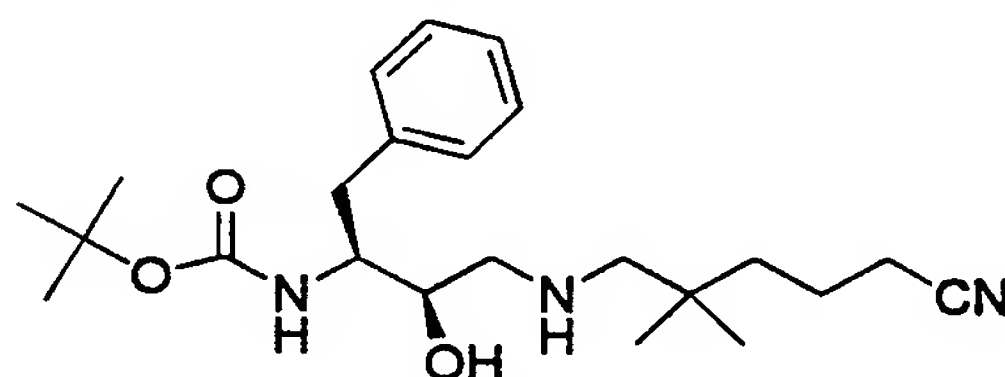
Step 1:

4-(1,3-Dioxolan-2-yl)-4-methylpentanenitrile

0.74 g (3.8 mmol) of 2-(4-chloro-1,1-dimethylbutyl)-1,3-dioxolane) and 0.50 g (7.7 mmol) of sodium cyanide in 5 mL of DMSO was heated to 100°C with stirring. After 18 hours the solution was cooled to RT, diluted with water, and the mixture extracted with ether (3x). The combined ether extracts were washed with water (3x), dried over MgSO₄, and concentrated *in vacuo* to afford 0.69 g (90%) of the desired compound as a light yellow liquid.

¹H NMR (CDCl₃) : δ 0.92 (6H, s), 1.41-1.48 (2H, m), 1.65-1.75 (2H, m), 2.32 (2H, t), 3.82-3.97 (4H, m), 4.53 (1H, s).

Step 2:

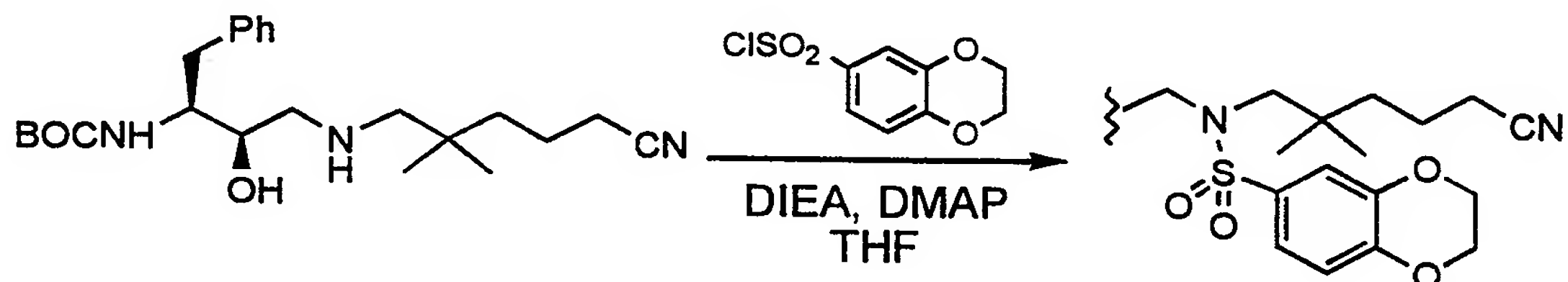
**tert-Butyl N-(1S,2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate**

The product from Step 1 (0.44 g (2.2 mmol) of 4-(1,3-dioxolan-2-yl)-4-methylpentyl azide) in 15 mL of THF was treated with 3 mL of 1M aqueous HCl and the solution stirred at reflux. After 4 hours the solution was cooled to RT and poured into 40 mL of rapidly stirred saturated aqueous NaHCO₃. The resulting mixture was extracted with ether (3x). The combined ether

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- extracts were washed with water (2x), dried over MgSO_4 and concentrated to a volume of approximately 3 mL. This solution was added to a solution of 0.62 g (2.2 mmol) of *tert*-butyl *N*-(1*S*,2*R*)-3-amino-1-benzyl-2-hydroxypropylcarbamate in 20 mL of 1:1 THF/DMF. The solution was treated with 1 g of powdered 4A molecular sieves and the mixture heated at 50°C. After 2 hours the mixture was cooled in an ice water bath and treated with 0.47 g (2.2 mmol) of $\text{NaBH}(\text{OAc})_3$. After stirring at RT for 18 hours the mixture was concentrated *in vacuo*. The residue was suspended in CH_2Cl_2 and stirred vigorously with 0.5M aqueous NaOH for 5 minutes. The layers were separated and the aqueous phase extracted with two additional portions of CH_2Cl_2 . The combined CH_2Cl_2 solutions were washed with water (2x), dried over MgSO_4 , and concentrated to give a clear viscous oil. This material was subjected to flash chromatography (SiO_2 , 95:5 CH_2Cl_2 /2M NH_3 in MeOH) to afford 0.45 g (49%) of the desired product as a white foam.
- ^1H NMR ($\text{DMSO}-d_6$): δ 0.82 (6H, s), 1.18-1.30 (11H, m), 1.40-1.53 (2H, m), 2.24 (2H, s), 2.38-2.62 (5H, m), 2.96 (1H, dd), 3.36-3.56 (2H, m), 4.75 (1H, br), 6.71 (1H, d), 7.08-7.26 (5H, m); MS 404 (MH+).

Step 3:

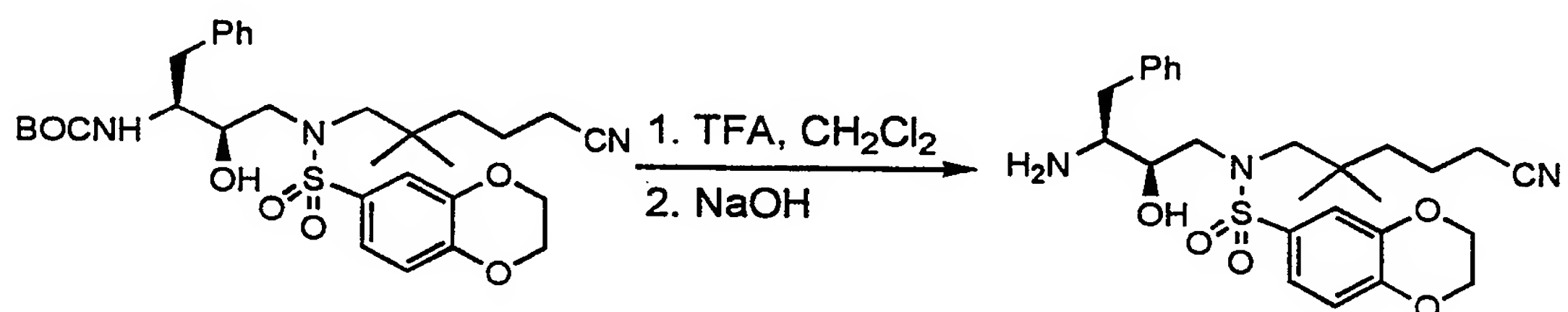


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tert-butyl *N*-(1*S*,2*R*)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)(2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl)amino]-2-hydroxypropylcarbamate

A solution of 0.35 g (0.87 mmol) of *tert*-butyl *N*-(1*S*,2*R*)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate, 0.22 g (0.95 mmol) of 3,4-ethylenedioxybenzenesulfonyl chloride, 0.18 mL (1.0 mmol) of *N,N*-diisopropylethylamine, and 10 mg of DMAP in 15 mL of anhydrous THF was stirred at RT. After 18 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 6:4 hexane/EtOAc) to afford 0.39 g (75%) of the desired product as a white foam. ¹H-NMR (CDCl₃): 7.32-7.17 (m, 7H), 6.92 (d, 1H), 4.57 (d, 1H), 4.30 (m, 4H), 4.11 (m, 1H), 3.90 (m, 1H), 3.67 (m, 1H), 3.18-3.02 (m, 2H), 3.01-2.90 (m, 2H), 2.80 (m, 2H), 2.31 (t, 2H), 1.70-1.53 (m, 2H), 1.49-1.19 (m, 2H), 1.25 (s, 9H), 0.92 (s, 6H). MS (ESI): 624 (M+Na).

Step 4:



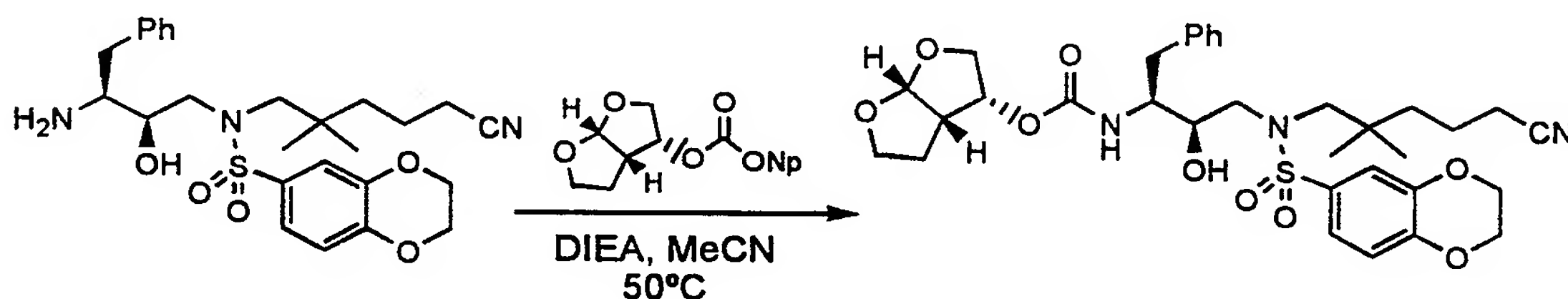
N-[(2*R*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl]-*N*-(5-cyano-2,2-dimethylpentyl)-2,3-dihydro-1,4-benzodioxine-6-sulfonamide.

A solution of 0.39 g (0.65 mmol) of *tert*-butyl *N*-(1*S*,2*R*)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)(2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl)amino]-2-hydroxypropylcarbamate in 25 mL of 1:1 TFA/CH₂Cl₂ was stirred at RT for 2 hours and was then concentrated *in*

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vacuo. The residue was dissolved in CH_2Cl_2 . The resulting solution was washed with 1M aqueous NaOH (1x), water (2x), dried over MgSO_4 , and concentrated to give 0.31 g (95%) of the desired amine as a white foam. $^1\text{H-NMR}$ (CDCl_3): 7.34–7.15 (m, 7H), 6.94 (d, 1H), 4.29 (m, 4H), 3.89 (m, 1H), 3.32–3.19 (m, 2H), 3.02 (m, 3H), 2.87 (dd, 1H), 2.41 (dd, 1H), 2.32 (t, 2H), 1.80–1.30 (7H), 0.96 (s, 6H). MS (ESI): 502 (M+H).

Step 5:



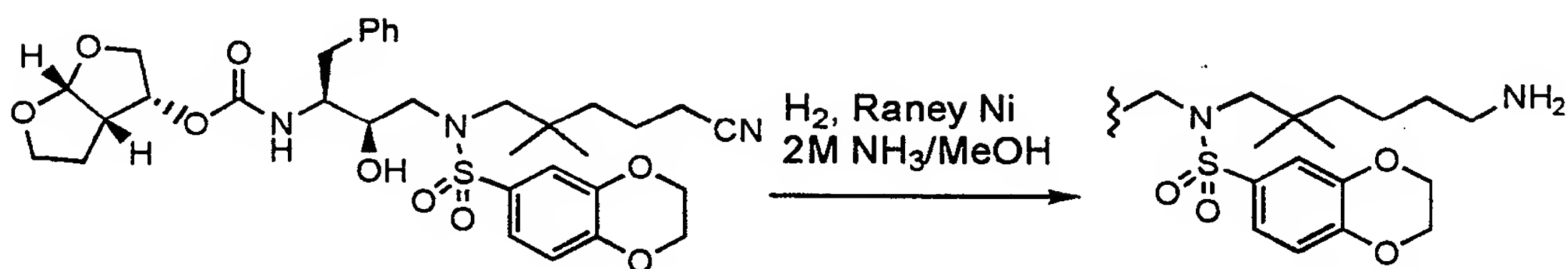
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)(2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl)amino]-2-hydroxypropyl]carbamate.

A solution of 0.31 g (0.62 mmol) of N-[(2R,3S)-3-amino-2-hydroxy-4-phenylbutyl]-N-(5-cyano-2,2-dimethylpentyl)-2,3-dihydro-1,4-benzodioxine-6-sulfonamide, 0.20 g (0.68 mmol) of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (4-nitrophenyl)carbonate, and 0.22 mL (1.2 mmol) of N,N-diisopropylethylamine in 20 mL of MeCN was heated to 50°C with stirring. After 18 hours the solution was cooled to RT and concentrated in vacuo. The residue was dissolved in CH_2Cl_2 . The resulting solution was washed with 1M aqueous NaOH (1x), water (3x), dried over MgSO_4 , and concentrated to dryness. The crude product was purified by flash chromatography (SiO_2 , 95:5 $\text{CH}_2\text{Cl}_2/\text{MeOH}$) to afford 0.36 g (88%) of the desired compound as a white foam. $^1\text{H-NMR}$

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NMR (CDCl₃): 7.30-7.12 (m, 7H), 6.96 (d, 1H), 5.60 (d, 1H), 4.99 (m, 2H), 4.29 (m, 4H), 4.02 (m, 2H), 3.90 (dd, 1H), 3.80 (t, 2H), 3.63 (m, 2H), 3.21-3.01 (m, 3H), 2.94 (d, 1H), 2.84 (m, 1H), 2.80-2.63 (m, 2H), 2.35 (t, 2H),
 5 1.70-1.30 (m, 6H), 0.94 (s, 3H), 0.90 (s, 3H). MS(ESI): 680 (M+Na).

Step 6:



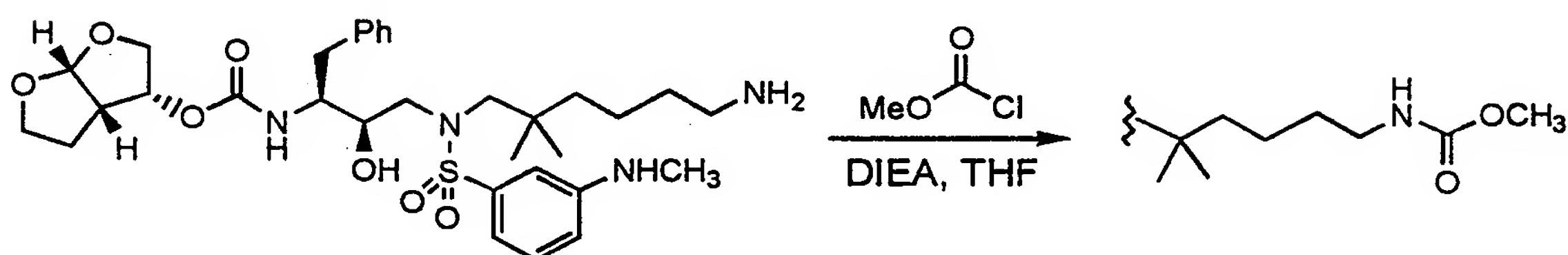
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(6-amino-2,2-dimethylhexyl) (2,3-dihydro-1,4-benzodioxin-
 10 6-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate

A mixture of 0.35 g (0.53 mmol) of
 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-
 benzyl-3-[(5-cyano-2,2-dimethylpentyl) (2,3-dihydro-1,4-
 benzodioxin-6-ylsulfonyl)amino]-2-hydroxypropylcarbamate
 15 and approximately 50 mg of Raney nickel (washed with
 water and then MeOH) in 40 mL of 2M NH₃/MeOH was subjected
 to hydrogenation at 45 psi. After 3 hours the vessel was
 purged with nitrogen, catalyst removed by filtration
 through celite and the filtrate concentrated to dryness
 20 in vacuo. The residue was subjected to flash
 chromatography (SiO₂, 9:1 CH₂Cl₂, 2M NH₃ in MeOH) to
 afford 0.31 g (89%) of the desired amine as a white foam.
 H¹-NMR (CDCl₃): 7.30-7.11 (m, 7H), 6.95 (d, 1H), 5.60 (d,
 1H), 5.28 (m, 1H), 4.99 (q, 1H), 4.30 (m, 4H), 4.00 (t,
 25 1H), 3.91 (dd, 1H), 3.81 (m, 2H), 3.66 (m, 2H), 3.17 (dd,
 1H), 3.10-2.97 (m, 3H), 2.83 (m, 2H), 2.79-2.62 (m, 3H),

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1.58 (m, 1H), 1.42-1.20 (m, 7H), 0.90 (d, 6H). MS (ESI): 662 (M+H).

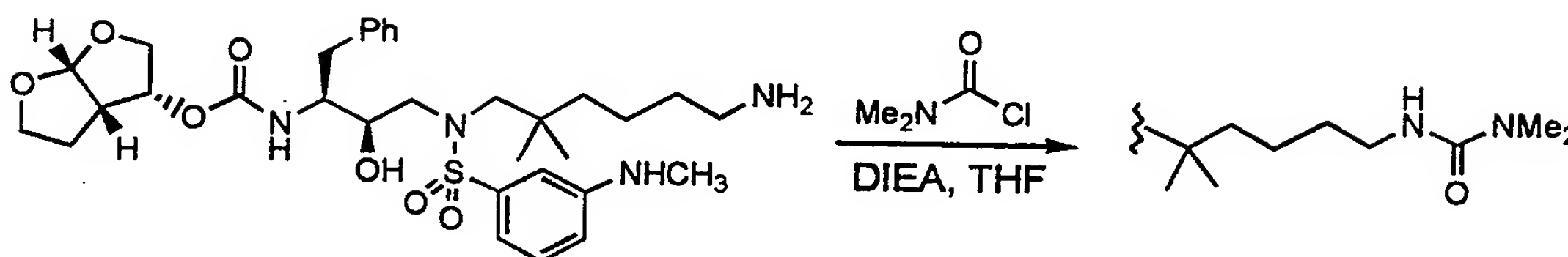
Example (Compound 246)



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-benzyl-2-hydroxy-3-(6-[(methoxycarbonyl)amino]-2,2-dimethylhexyl[3-(methylamino)phenyl]sulfonylamino)propyl]carbamate.

A solution of 55 mg (0.087 mmol) of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-((6-amino-2,2-dimethylhexyl)[3-(methylamino)phenyl]sulfonylamino)-1-benzyl-2-hydroxypropyl]carbamate and 17.0 μ L (0.091 mmol) of N,N-diisopropylethylamine in 4 mL of anhydrous THF at 0°C was treated with 7.0 μ L (0.091 mmol) of methyl chloroformate. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 55 mg (92%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.34-7.12 (m, 6H), 7.02 (d, 1H), 6.94 (s, 1H), 6.78 (d, 1H), 5.60 (d, 1H), 5.17 (d, 1H), 5.02-4.70 (m, 2H), 4.14 (s, 1H), 4.02 (m, 1H), 3.89 (dd, 1H), 3.79 (m, 2H), 3.72-3.55 (m, 5H), 3.28-2.91 (m, 6H), 2.83 (s, 3H), 2.89-2.64 (m, 3H), 1.65-1.18 (m, 8H), 0.91 (s, 3H), 0.88 (s, 3H). MS (ESI): 691 (M+H).

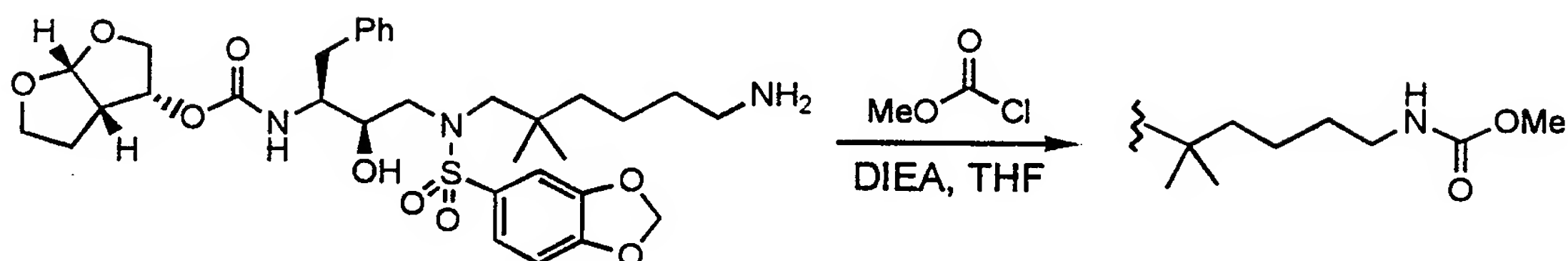
-226-

Example (Compound 247)

(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*,2*R*)-1-benzyl-3-((6-[(dimethylamino) carbonyl] amino-2,2-dimethylhexyl) [3-(methylamino) phenyl] sulfonylamino)-2-hydroxypropyl] carbamate.

A solution of 55 mg (0.087 mmol) of (3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*,2*R*)-3-((6-amino-2,2-dimethylhexyl) [3-(methylamino) phenyl] sulfonylamino)-1-benzyl-2-hydroxypropyl] carbamate and 17 μ L (0.091 mmol) of *N,N*-diisopropylethylamine in 4 mL of anhydrous THF at 0°C was treated with 8.4 μ L (0.091 mmol) of *N,N*-dimethylcarbamyl chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated in vacuo and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 49 mg (80%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.32-7.12 (m, 6H), 7.02 (d, 1H), 6.92 (s, 1H), 6.76 (d, 1H), 5.60 (m, 2H), 4.96 (q, 1H), 4.51 (m, 1H), 4.32 (s, 1H), 4.20 (br s, 1H), 4.01 (m, 1H), 3.88 (dd, 1H), 3.79 (m, 2H), 3.62 (m, 2H), 3.30-2.92 (m, 6H), 2.91-2.64 (m, 12H), 1.65-1.40 (m, 3H), 1.38-1.20 (m, 5H), 0.91 (s, 3H), 0.85 (s, 3H). MS (ESI): 704 (M+H).

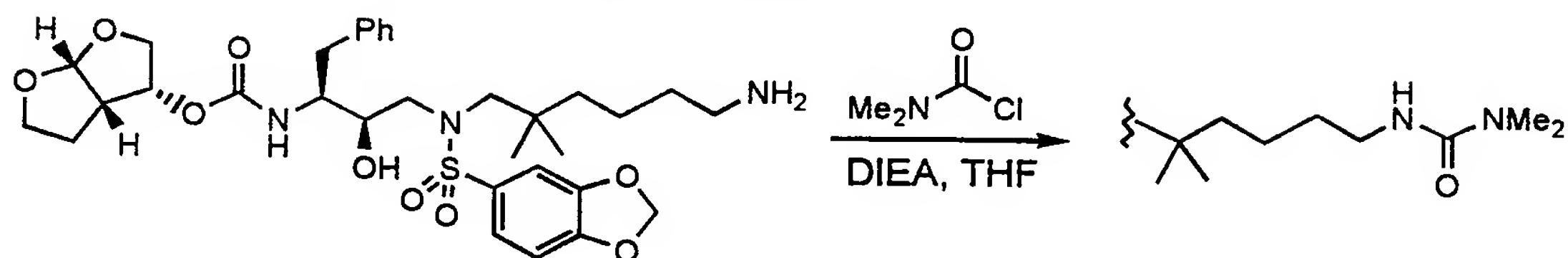
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Example (Compound 248)

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-
 ((1,3-benzodioxol-5-ylsulfonyl)6-
 [(methoxycarbonyl)amino]-2,2-dimethylhexylamino)-1-
 5 benzyl-2-hydroxypropyl]carbamate.

A solution of 60 mg (0.093 mmol) of
 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(6-amino-2,2-dimethylhexyl)(1,3-benzodioxol-5-
 ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate and
 10 20 μ L (0.11 mmol) of N,N-diisopropylethylamine in 5 mL of
 anhydrous THF at 0°C was treated with 8.0 μ L (0.10 mmol)
 of methyl chloroformate. The solution was allowed to
 warm to RT with stirring. After 18 hours the solution
 was concentrated *in vacuo* and the residue was subjected
 15 to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH)
 to afford 59 mg (89%) of the desired compound as a white
 foam. ¹H-NMR (CDCl₃): 7.40 (dd, 1H), 7.35-7.20 (m, 6H),
 6.96 (d, 1H), 6.14 (s, 2H), 5.65 (d, 1H), 5.27 (d, 1H),
 5.12-4.90 (m, 2H), 4.20 (br s, 1H), 4.08 (m, 1H), 3.98
 20 (dd, 1H), 3.86 (m, 2H), 3.80-3.63 (m, 5H), 3.30-2.71 (m,
 9H), 1.62-1.23 (m, 8H), 0.96 (s, 3H), 0.92 (s, 3H).
 MS (ESI): 728 (M+Na).

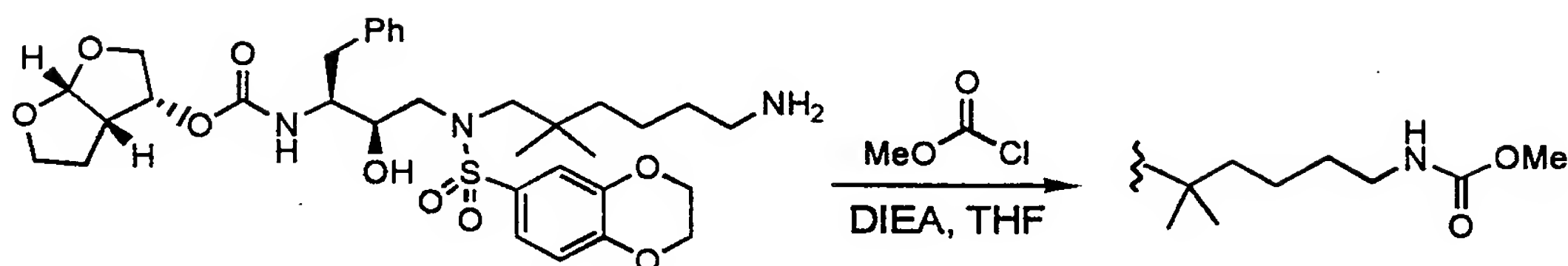
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Example (Compound 249)

(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(6-
 5 [(dimethylamino) carbonyl]amino-2,2-dimethylhexyl) amino]-
 1-benzyl-2-hydroxypropylcarbamate.

A solution of 60 mg (0.093 mmol) of
 (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-3-
 [(6-amino-2,2-dimethylhexyl)(1,3-benzodioxol-5-
 10 ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate and
 20 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 5 mL of
 anhydrous THF at 0°C was treated with 9.4 μ L (0.10 mmol)
 of *N,N*-dimethylcarbonyl chloride. The solution was
 allowed to warm to RT with stirring. After 18 hours the
 15 solution was concentrated *in vacuo* and the residue was
 subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃
 in MeOH) to afford 60 mg (90%) of the desired compound as
 a white foam. ¹H-NMR (CDCl₃): 7.39 (dd, 1H), 7.32-7.16 (m,
 6H), 6.96 (d, 1H), 6.12 (s, 2H), 5.78 (d, 1H), 5.63 (d,
 20 1H), 5.02 (q, 1H), 4.58 (t, 1H), 4.45 (br s, 1H), 4.07
 (m, 1H), 3.95 (dd, 1H), 3.82 (m, 2H), 3.68 (m, 2H), 3.32-
 3.01 (m, 6H), 2.91 (s, 6H), 2.90-2.69 (m, 3H), 1.70-1.43
 (m, 3H), 1.42-1.25 (m, 5H), 1.00 (s, 3H), 0.91 (s, 3H).
 MS (ESI): 719 (M+H).

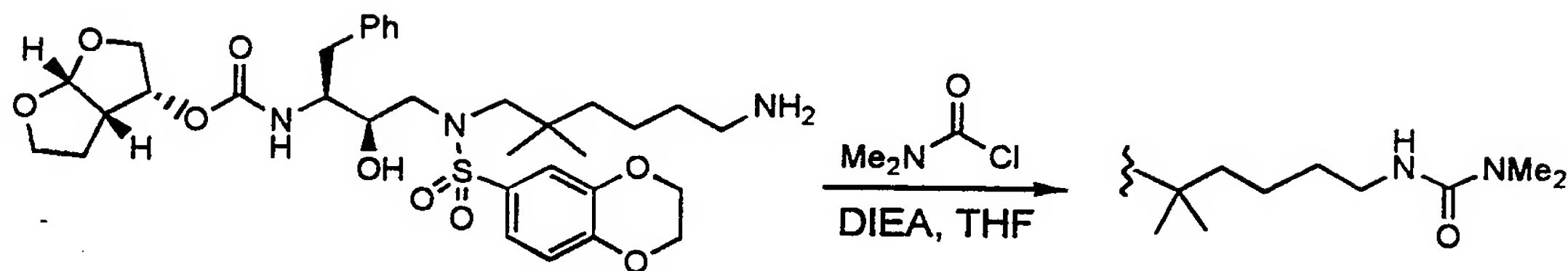
-229-

Example (Compound 250)

(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*, 2*R*)-1-benzyl-3-((2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) 6-[(methoxycarbonyl) amino]-2,2-dimethylhexylamino)-2-hydroxypropyl] carbamate.

A solution of 60 mg (0.091 mmol) of (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-3-[(6-amino-2,2-dimethylhexyl) (2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate and 20 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 7.7 μ L (0.10 mmol) of methyl chloroformate. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 65 mg (98%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.31-7.13 (m, 7H), 6.94 (d, 1H), 5.60 (d, 1H), 5.21 (d, 1H), 5.02-4.83 (m, 2H), 4.30 (m, 4H), 4.19 (br s, 1H), 4.01 (m, 1H), 3.89 (dd, 1H), 3.80 (m, 2H), 3.72-3.59 (m, 5H), 3.22-3.03 (m, 5H), 2.93 (d, 1H), 2.83 (m, 1H), 2.73 (m, 2H), 1.60-1.41 (m, 3H), 1.40-1.18 (m, 5H), 0.91 (s, 3H), 0.89 (s, 3H). MS (ESI): 742 (M+Na).

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Example (Compound 251)

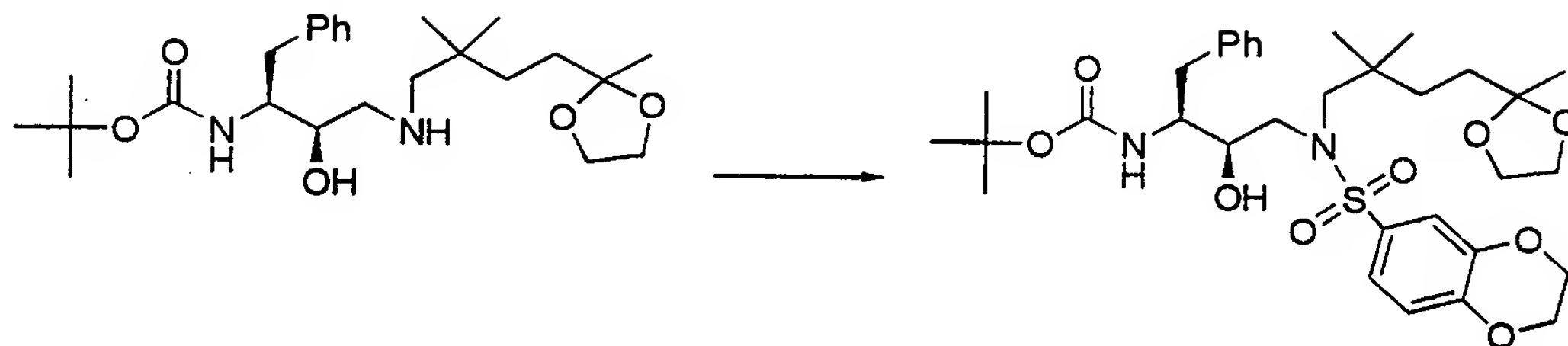
(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-1-benzyl-3-[(2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) (6-[(dimethylamino) carbonyl] amino-2,2-dimethylhexyl) amino]-2-hydroxypropylcarbamate.

A solution of 60 mg (0.091 mmol) of (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-3-[(6-amino-2,2-dimethylhexyl) (2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate and 20 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 9.2 μ L (0.10 mmol) of *N,N*-dimethylcarbamyl chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 54 mg (81%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.31-7.11 (m, 7H), 6.96 (d, 1H), 5.67 (d, 1H), 5.60 (d, 1H), 4.96 (q, 1H), 4.51 (m, 1H), 4.43 (br s, 1H), 4.30 (m, 4H), 4.01 (t, 1H), 3.90 (dd, 1H), 3.78 (m, 2H), 3.63 (m, 2H), 3.24-2.92 (m, 6H), 2.87 (s, 6H), 2.87-2.67 (m, 3H), 1.60-1.40 (m, 3H), 1.39-1.20 (m, 5H), 0.92 (s, 3H), 0.86 (s, 3H). MS (ESI): 733 (M+H).

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Example (Compound 252)

Step 1:

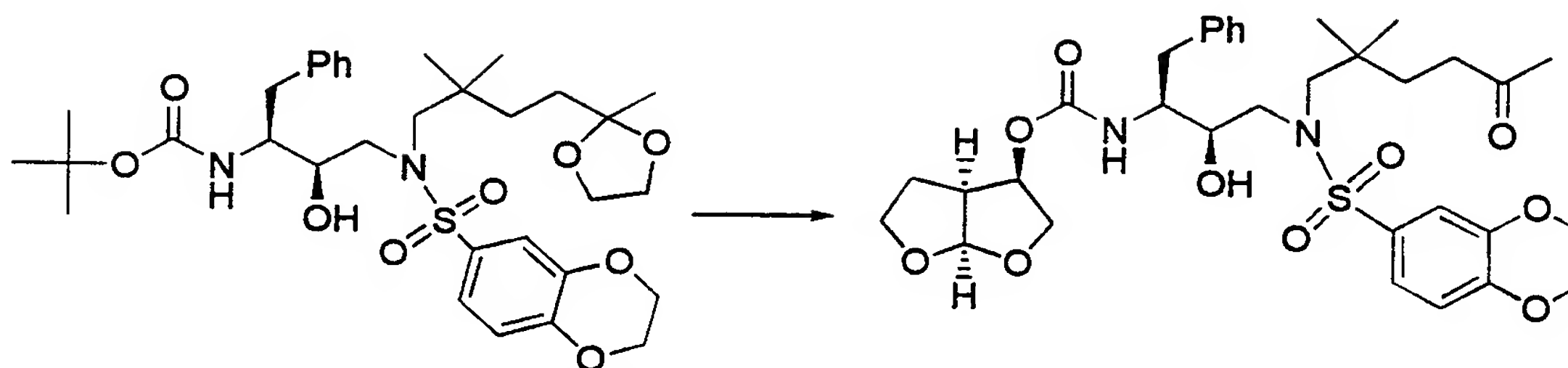


tert-Butyl N-((1S,2R)-3-(1,4-benzodioxan-6-sulfonyl)[2,2-dimethyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl]amino-1-benzyl-2-hydroxypropyl)carbamate.

The product from step 2 (Example (Compound 208)) was reacted with 1,4-benzodioxan-6-sulfonyl chloride according to procedure 3 (Example (Compound 201)) to give the title compound as a foam; ^1H NMR (DMSO- d_6): δ 0.9 (6H, s), 1.1 (1H, s), 1.2 (9H, s), 1.3 (2H, t), 1.5 (2H, t), 2.45-2.50 (1H, m), 2.8 (1H, d), 2.9 (1H, dd), 3.0 (1H, d), 3.2-3.4 (5H, m), 3.75 (1H, quartet), 3.80-3.85 (4H, m), 4.25-4.35 (4H, m), 5.0 (1H, d), 6.6 (1H, d), 7.0 (1H, d), 7.1-7.3 (7H, m); MS: 671.2 (MH^+); $\text{C}_{33}\text{H}_{48}\text{N}_2\text{O}_9\text{S}$.

Step 2:

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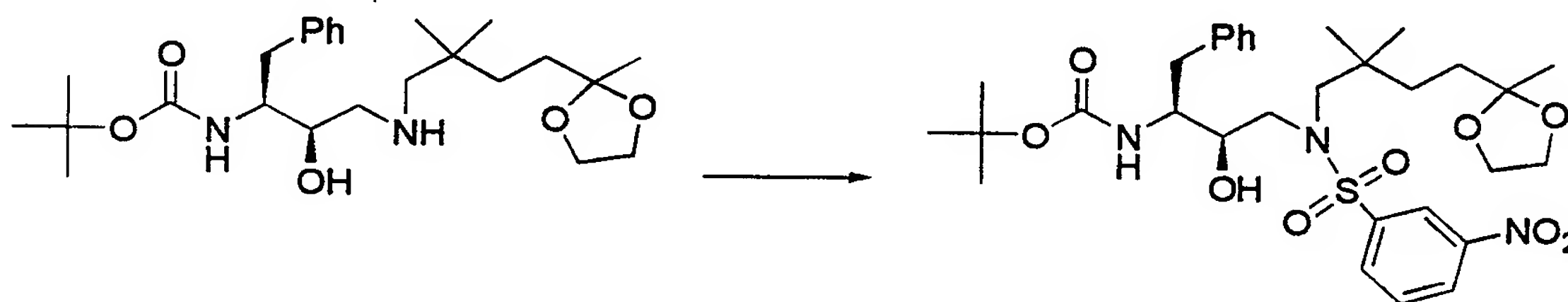


(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-(1,4-benzodioxan-6-sulfonyl)-2-hydroxypropyl)-1-benzyl-2-hydroxypropyl carbamate

The product from step 1 was subjected to the procedure used in Example (Compound 205)) to afford the title compound as a foam; ^1H NMR ($\text{DMSO}-d_6$): δ 0.86 (3H, s), 0.90 (3H, s), 1.1-1.2 (1H, m), 1.25-1.35 (1H, m), 1.45 (2H, t), 2.1 (3H, s), 2.35-2.45 (3H, m), 2.7-2.8 (2H, m), 2.82-3.00 (2H, m), 3.25-3.30 (2H, m), 3.4 (1H, br quartet), 3.5-3.6 (2H, m), 3.7 (1H, d), 3.75-3.85 (2H, m), 4.25-4.35 (4H, m), 4.8 (1H, dt), 5.18 (1H, d), 5.5 (1H, d), 7.0 (1H, d), 7.1-7.3 (8H, m); MS: 661.2 (MH^+); $\text{C}_{33}\text{H}_{44}\text{N}_2\text{O}_{10}\text{S}$.

Example (Compound 253)

Step 1:



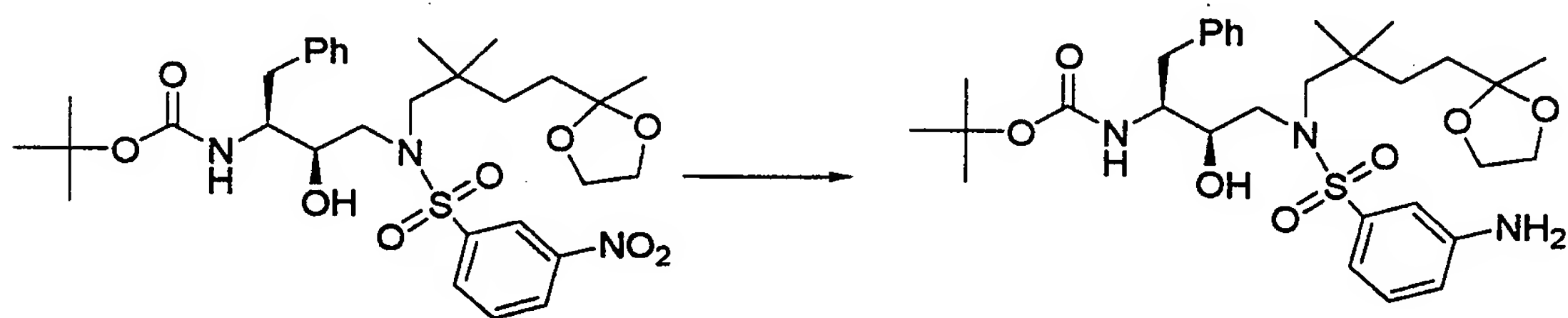
tert-Butyl N-((1S,2R)-1-benzyl-3-([2,2-dimethyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl])butyl)-2-hydroxypropyl carbamate-3-(3-nitrophenyl)sulfonyl

The product from step 2 (Example (Compound 208)) was reacted with 3-nitrobenzenesulfonyl chloride

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according to procedure 3 (Example (Compound 201)) to give the title compound as a foam; ^1H NMR ($\text{DMSO}-d_6$): δ 0.9 (3H, s), 0.93 (3H, s), 1.1 (1H, s), 1.15 (9H, s), 1.3-1.4 (2H, m), 1.5-1.6 (2H, m), 2.4 (1H, dd), 2.8 (1H, d), 3.0 (1H, d), 3.2-3.4 (4H, m), 3.4-3.5 (3H, m), 3.8 (4H, br s), 4.85 (1H, d), 6.6 (1H, d), 7.1-7.3 (5H, m), 7.85 (1H, t), 8.22 (1H, d), 8.45-8.55 (2H, m).

Step 2:

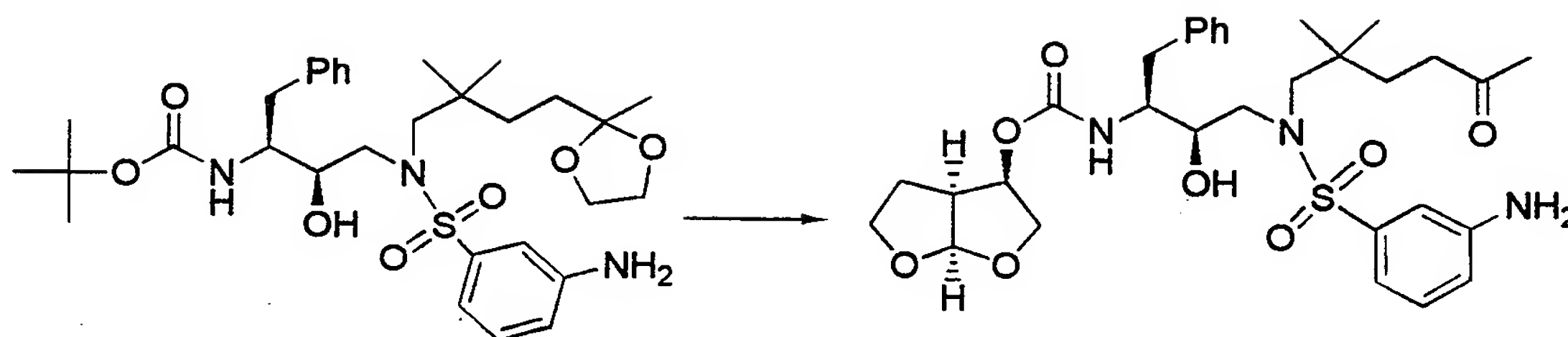


***tert*-Butyl *N*-((1*S*,2*R*)-3-([(3-aminophenyl)sulfonyl][2,2-dimethyl-4-(2-methyl-1,3-dioxolan-2-yl)butyl]amino-1-benzyl-2-hydroxypropyl)carbamate**

A mixture of the product from step 1 (1.92 g, 3.0 mmol), 10% palladium/carbon (500 mg), and 2M ammonia/ethanol (40 mL) was hydrogenated at atmospheric pressure for 4.5 hours. The mixture was diluted with dichloromethane and filtered through celite. Solvent was evaporated and dried in vacuo to provide the title compound (1.6 g, 87%) as a white foam; ^1H NMR ($\text{DMSO}-d_6$): δ 0.9 (6H, s), 1.1 (1H, s), 1.2 (9H, s), 1.3 (2H, t), 11.5 (2H, t), 2.4-2.5 (1H, m), 2.8 (1H, d), 2.9-3.0 (2H, m), 3.2-3.5 (5H, m), 3.75-3.80 (1H, m), 3.82 (4H, s), 4.95 (1H, d), 5.55 (2H, s), 6.58 (1H, d), 6.75 (1H, d), 6.85 (1H, d), 7.0 (1H, s), 7.1-7.3 (6H, m).

Step 3:

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(3*R*, 3*aS*, 6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-3-
 ([(3-aminophenyl) sulfonyl] (2,2-dimethyl-5-oxohexyl) amino-
 1-benzyl-2-hydroxypropyl) carbamate

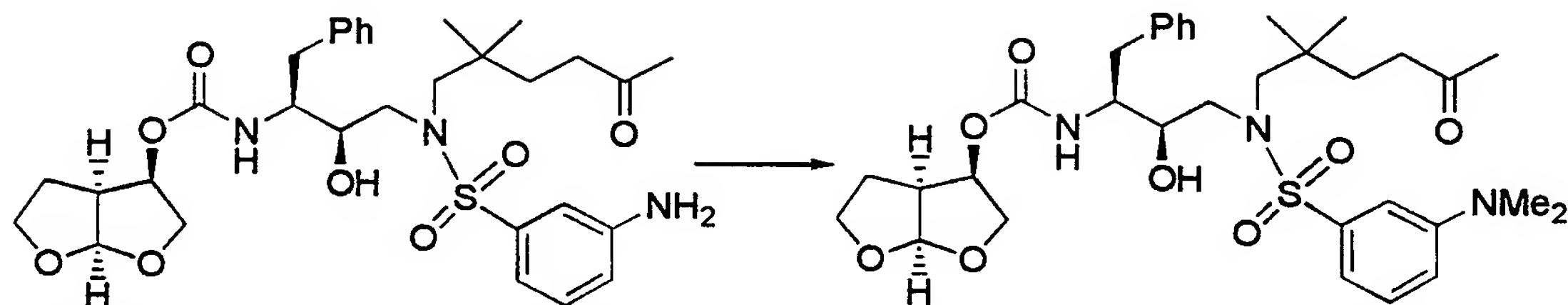
The product from step 2 was subjected to the
 5 procedure used in Example (Compound 205) to afford the
 title compound as a foam; ¹H NMR (DMSO-*d*₆): δ 0.85 (3H,
 s), 0.90 (3H, s), 1.1-1.2 (1H, m), 1.3-1.4 (1H, m), 1.45
 (2H, t), 2.1 (3H, s), 2.25-2.35 (3H, m), 2.7-2.8 (2H, m),
 2.85 (1H, dd), 2.90 (1H, dd), 3.30-3.35 (2H, m), 3.45
 10 (1H, br quartet), 3.55 (1H, dd), 3.58-3.60 (1H, m), 3.7
 (1H, td), 3.75-3.85 (2H, m), 4.8 (1H, dt), 5.1 (1H, d),
 5.48 (1H, d), 5.55 (2H, br s), 6.77 (1H, dd), 6.85 (1H,
 br d), 6.97 (1H, s), 7.10-7.25 (7H, m); MS: 618.2 (MH⁺);
 C₃₁H₄₃N₃O₈S.

15

Example (Compound 254)

Step 1:

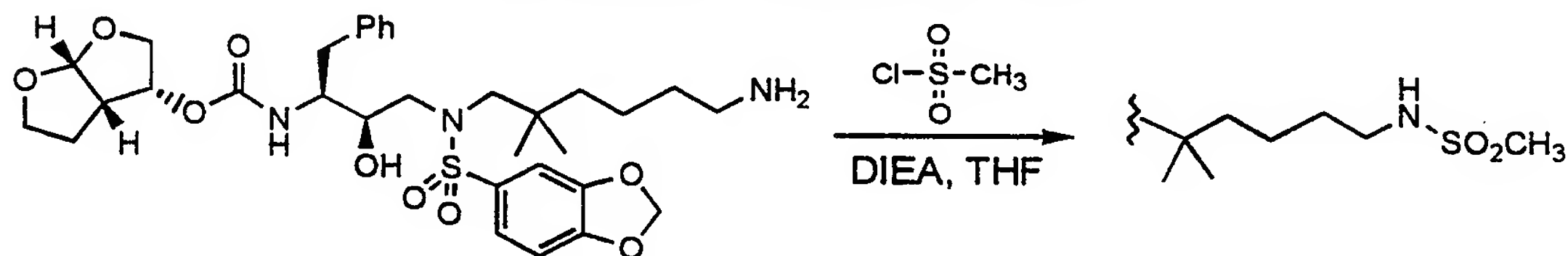
-235-



(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-
 ([(3-dimethylaminophenyl) sulfonyl] (2,2-dimethyl-5-
 oxohexyl) amino-1-benzyl-2-hydroxypropyl) carbamate

The product from step 1 (Example (Compound
 5 253)) was dissolved (0.10g) in ethanol (3 mL) and 37%
 HCOH/water (0.4 mL) was hydrogenated at atmospheric
 pressure with 10% Pd/C (25 mg) for 18 hours. The mixture
 was filtered through a bed celite, evaporated, and
 dissolved in dichloromethane. The solution was washed
 10 with water, dried (sodium sulfate), evaporated, and dried
 in vacuo to provide the title compound (90 mg) as a foam.
 0.87 (3H, s), 0.88 (3H, s), 1.1-1.2 (1H, m), 1.25-1.40
 (1H, m), 1.45 (2H, t), 2.05 (3H, s), 2.30-2.45 (3H, m),
 2.7-2.8 (2H, m), 2.9 (8H, br s), 3.2-3.3 (2H, m), 3.4-3.6
 15 (3H, m), 3.7 (1H, t), 3.8 (1H, d), 3.85 (1H, d), 4.8 (1H,
 dt), 5.1 (1H, d), 5.5 (1H, d), 6.97 (1H, d), 6.98 (1H,
 s), 7.0 (1H, d), 7.10-7.22 (6H, m), 7.35 (1H, t); MS:
 646.2 (MH⁺); C₃₃H₄₇N₃O₈S.

Example (Compound 255)



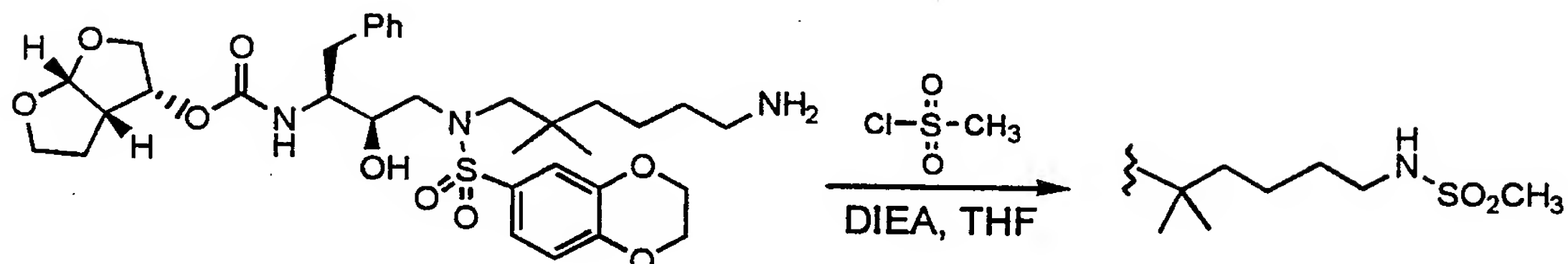
20

(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-
 ((1,3-benzodioxol-5-ylsulfonyl)2,2-dimethyl-6-
 [(methylsulfonyl)amino]hexylamino)-1-benzyl-2-
 hydroxypropyl]carbamate

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A solution of 60 mg (0.093 mmol) of (3*R*, 3*aS*, 6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-3-[(6-amino-2,2-dimethylhexyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate and 5 20 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 7.9 μ L (0.10 mmol) of methanesulfonyl chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected 10 to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 58 mg (85%) of the desired compound as a white foam. H¹-NMR (CDCl₃): 7.39 (dd, 1H), 7.34-7.20 (m, 6H), 6.96 (d, 1H), 6.15 (s, 2H), 5.66 (d, 1H), 5.03 (m, 1H), 4.82 (t, 1H), 4.16-3.81 (m, 5H), 3.72 (m, 2H), 3.26-2.72 15 (m, 9H), 2.99 (s, 3H), 1.60 (m, 3H), 1.49-1.30 (m, 5H), 0.99 (m, 6H). MS (ESI): 748 (M+Na).

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Example (Compound 256)

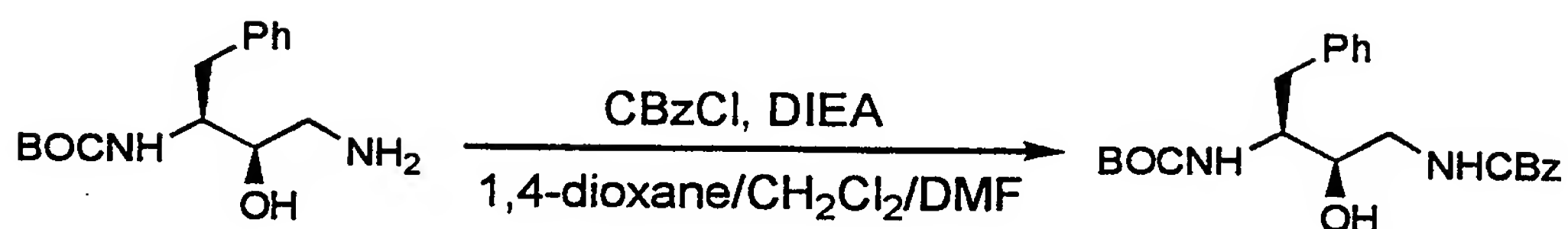
(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*,2*R*)-1-benzyl-3-((2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl)2,2-dimethyl-6-[(methanesulfonyl)amino]hexylamino)-2-hydroxypropyl] carbamate.

A solution of 60 mg (0.091 mmol) of (3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-3-[(6-amino-2,2-dimethylhexyl)(2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate and 20 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 7.8 μ L (0.10 mmol) of methanesulfonyl chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 58 mg (87%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.37-7.19 (m, 7H), 7.02 (d, 1H), 5.67 (d, 1H), 5.02 (m, 2H), 4.85 (t, 1H), 4.36 (m, 4H), 4.10 (m, 2H), 4.00-3.81 (m, 3H), 3.80-3.64 (m, 2H), 3.25-2.70 (m, 9H), 2.99 (s, 3H), 1.60 (m, 3H), 1.49-1.23 (m, 5H), 0.99 (m, 6H). MS (ESI): 762 (M+Na).

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Example (Compound 257)

Step 1:

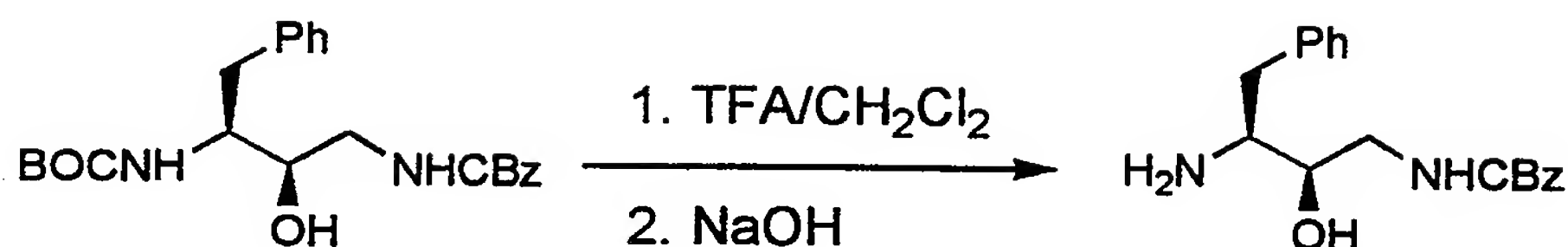


Benzyl *N*-[(2*R*,3*S*)-3-[(*tert*-butoxycarbonyl)amino]-2-hydroxy-4-phenylbutyl]carbamate

5 A solution of 20.0 g (71.3 mmol) of *tert*-butyl *N*-[(1*S*,2*R*)-3-amino-1-benzyl-2-hydroxypropyl]carbamate in 1 L of 4:1:1 dioxane/CH₂Cl₂/DMF at 0°C was treated with 13.7 mL (78.4 mmol) of *N,N*-diisopropylethylamine. A small amount of solid precipitated at this point. The mixture was treated with 10.2 mL (71.3 mmol) of benzyl chloroformate by slow addition over 5 minutes and then allowed to warm to RT. A clear solution resulted which was stirred at RT. After 18 hours the solution was concentrated *in vacuo*. The residue was suspended (would not dissolve) in CH₂Cl₂ and the mixture was shaken with 10% aqueous citric acid in a separatory funnel. The aqueous suspension was separated, filtered to remove solid and the aqueous filtrate discarded. The solid was combined with the CH₂Cl₂ suspension and the mixture concentrated to dryness at reduced pressure. The resulting residue was recrystallized from EtOH/water to afford 24.1 g (81%) of the desired compound at a white crystalline solid. NMR (DMSO-*d*₆): 7.43-7.03 (m, 11H), 6.63 (d, 1H), 5.04 (s, 2H), 4.99 (d, 1H), 3.61-3.40 (m, 2H), 3.30 (m, 1H), 3.06-2.84 (m, 2H), 2.52 (m, 1H), 1.27 (s, 9H). MS(ESI): 437 (M+Na).

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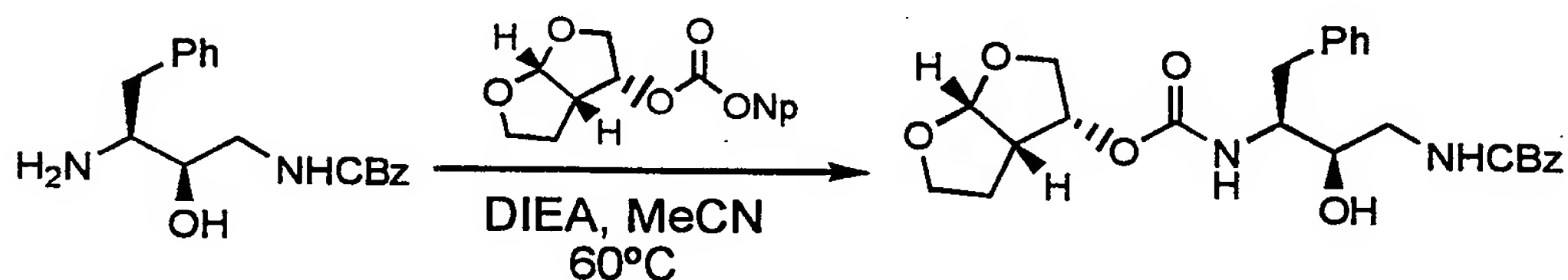
Step 2:



Benzyl N-[(2*R*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl]carbamate

A solution of 7.0 g (16.9 mmol) of benzyl N-(2*R*,3*S*)-3-[(*tert*-butoxycarbonyl)amino]-2-hydroxy-4-phenylbutylcarbamate in 100 mL of 1:1 TFA/CH₂Cl₂ was stirred at RT. After 2.5 hours the solution was concentrated *in vacuo*. The residue was dissolved in 1:1 CH₂Cl₂/THF and the solution washed with 1M aqueous NaOH (1x) followed by water (2x). The solution was not exposed to drying agent as this induced crystallization but was instead directly concentrated to dryness at reduced pressure. The residual water was removed by storing in *vacuo* overnight. This afforded the desired amine as a white solid in quantitative yield. NMR (DMSO-*d*₆): 7.38–7.06 (m, 11H), 5.00 (s, 2H), 4.73 (d, 1H), 3.25 (m, 2H), 3.04 (m, 1H), 2.83 (dd, 1H), 2.71 (m, 1H), 2.29 (dd, 1H), 1.15 (br s, 2H). MS(ESI): 315(M+H).

Step 3:

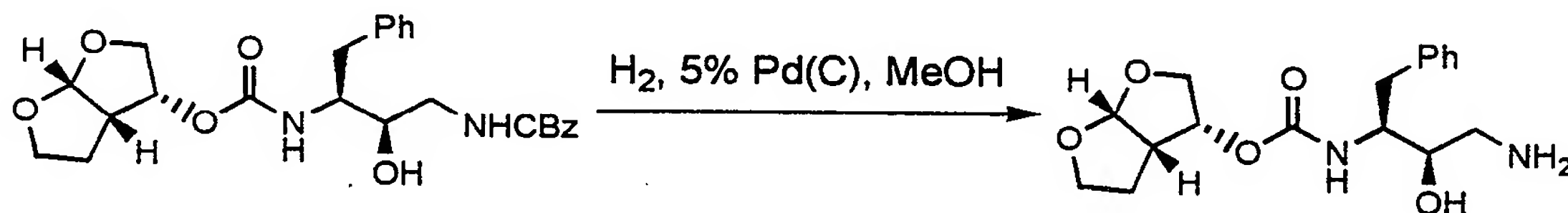


3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl N-((1*S*,2*R*)-1-benzyloxy) carbonyl]amino-2-hydroxypropyl) carbamate.

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A solution of 5.31 g (16.9 mmol) of benzyl N-[(2*R*,3*S*)-3-amino-2-hydroxy-4-phenylbutyl]carbamate, 4.99 g (16.9 mmol) of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl (4-nitrophenyl)carbonate and 5.9 mL (33 mmol) of *N,N*-diisopropylethylamine in 250 mL of MeCN was heated to 60°C with stirring under nitrogen. After 24 hours product had precipitated from the solution giving a yellow suspension. The mixture was cooled to RT at which point additional product precipitated. The solid was collected by filtration and dried *in vacuo* to give 4.9 g of material. The filtrate was concentrated to dryness and the residue recrystallized from MeCN to afford an additional 0.85 g of product for a total yield of 5.8 g (73%). NMR (DMSO-*d*₆): 7.42–7.09 (m, 12H), 5.51 (d, 1H), 5.06 (m, 3H), 4.84 (q, 1H), 3.83 (dd, 1H), 3.72 (t, 1H), 3.67–3.41 (m, 4H), 3.28 (m, 1H), 3.10–2.72 (m, 4H), 2.41 (m, 1H), 1.26 (m, 1H). MS(ESI): 493(M+Na).

Step 4:



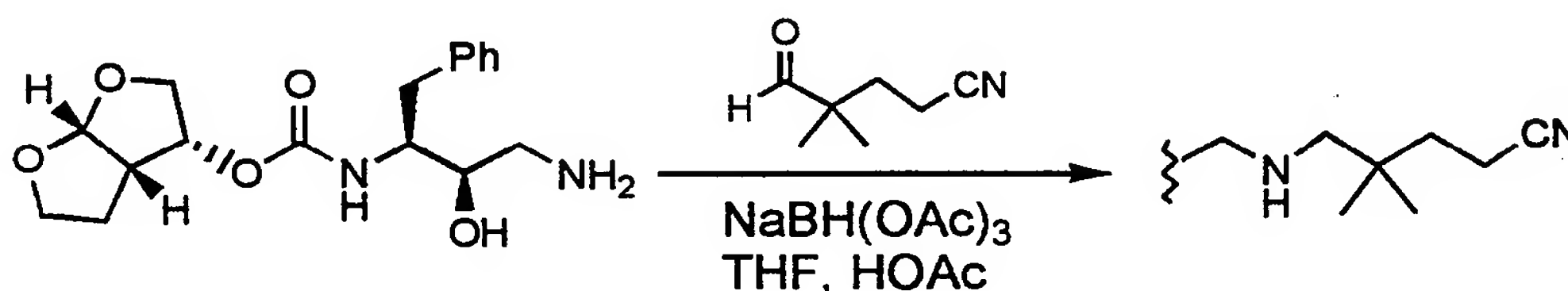
(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl N-[(1*S*,2*R*)-3-amino-1-benzyl-2-hydroxypropyl]carbamate.

A suspension of 4.0 g (8.5 mmol) of (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl N-[(1*S*,2*R*)-1-benzyl-3-[(benzyloxy)carbonyl]amino-2-hydroxypropyl]carbamate and 0.5 g of 5% Pd(C) in 200 mL of MeOH was subjected to hydrogenation at 45 psi. After 3.5 hours the reaction vessel was purged with nitrogen, catalyst was removed by filtration through celite and the

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filtrate was concentrated *in vacuo* to afford 2.72 g (95%) of the desired product as a white solid. NMR (DMSO- d_6): 7.23-7.04 (m, 6H), 5.45 (d, 1H), 4.91-4.75 (m, 2H), 3.79 (dd, 1H), 3.68 (t, 1H), 3.53 (m, 3H), 3.21 (m, 2H), 3.01 (dd, 1H), 2.74 (q, 1H), 2.57 (dd, 1H), 2.41 (m, 1H), 1.60-1.28 (m, 3H), 1.22 (dd, 1H). MS(ESI): 337(M+H).

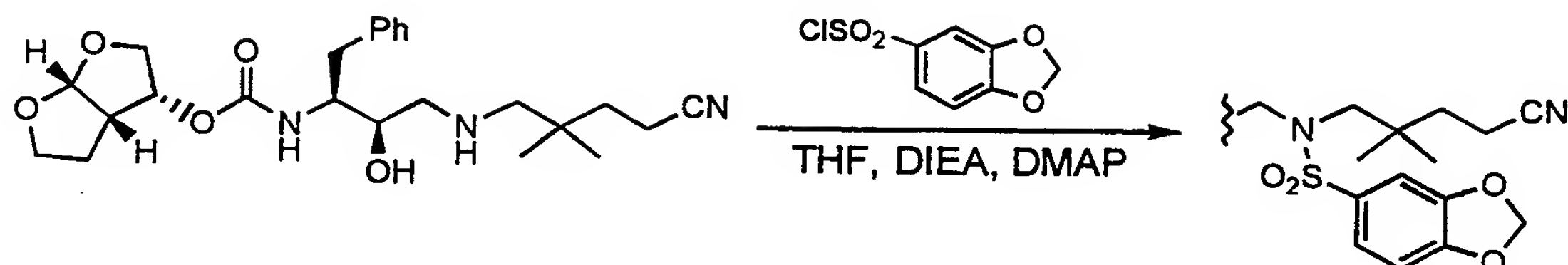
Step 5:



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-amino-1-benzyl-2-hydroxypropyl]carbamate and 1.32 g (10.5 mmol) of 4-cyano-2,2-dimethylbutyraldehyde in 175 mL of 3:1 THF/glacial HOAc was treated with 2.23 g (10.5 mmol) of NaBH(OAc)₃ and the resulting solution was stirred at RT. After 18 hours the solution was concentrated *in vacuo* and the residue dissolved in CH₂Cl₂. The solution was washed with 1M aqueous NaOH (1x), brine (2x), dried over MgSO₄, and concentrated to give a viscous oil. The crude product was purified by flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 3.2 g (75%) of the desired compound as a white foam. NMR (CDCl₃): 7.38-7.20 (m, 5H), 5.67 (d, 1H), 5.08 (m, 2H), 4.04-3.81 (m, 3H), 3.72 (m, 2H), 3.59 (q, 1H), 3.13 (dd, 1H), 2.96 (m, 1H), 2.79 (m, 3H), 2.42 (s, 2H), 2.32 (t, 2H), 1.79-1.43 (m, 6H), 1.00 (m, 6H). MS(ESI): 446(M+H).

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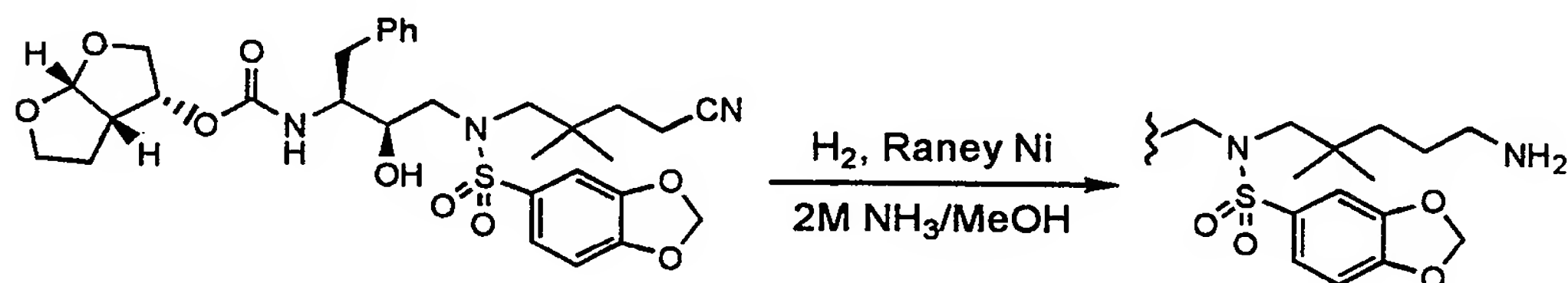
Step 6:



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(4-cyano-2,2-dimethylbutyl)amino]-1-benzyl-2-hydroxypropylcarbamate

- 5 A solution of 0.50 g (1.1 mmol) of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-cyano-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate, 0.27 g (1.2 mmol) of 3,4-methylenedioxybenzenesulfonyl chloride, 0.23 mL (1.3 mmol) of N,N-diisopropylethylamine, and 13 mg of DMAP in 15 mL of anhydrous THF was stirred at RT. After 18 hours the solution was concentrated in vacuo and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.63 g (91%) of the desired product as
- 15 a white foam. NMR (CDCl₃): 7.32 (d, 1H), 7.30-7.14 (m, 6H), 6.90 (d, 1H), 6.09 (s, 2H), 5.61 (d, 1H), 4.99 (q, 1H), 4.82 (d, 1H), 4.02 (m, 1H), 3.92 (dd, 1H), 3.81 (m, 3H), 3.76-3.58 (m, 3H), 3.14 (dd, 1H), 3.04 (m, 2H), 2.86 (m, 2H), 2.71 (dd, 1H), 2.37 (t, 2H), 1.74 (t, 2H), 1.58 (m, 1H), 1.39 (m, 1H), 0.98 (s, 6H). MS(ESI): 652 (M+Na).
- 20

Step 7:



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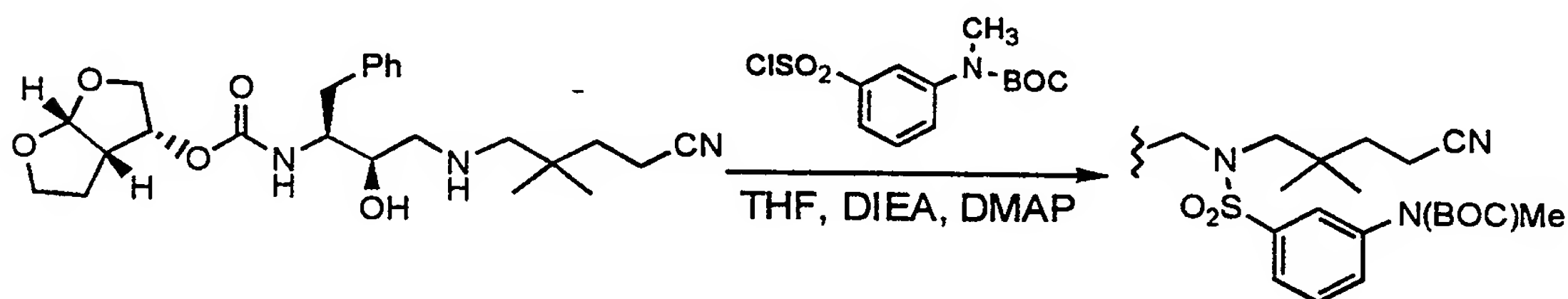
(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-3-[(5-amino-2,2-dimethylpentyl) (1,3-benzodioxol-5-ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate

A mixture of 0.63 g (0.99 mmol) of
 5 (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (4-cyano-2,2-dimethylbutyl) amino]-1-benzyl-2-hydroxypropylcarbamate and approximately 150 mg of Raney nickel (washed with water and MeOH) in 35 mL of 2M NH₃/MeOH was subjected to
 10 hydrogenation at 45 psi. After 3 hours the reaction vessel was purged with nitrogen, catalyst removed by filtration through celite and the filtrate concentrated *in vacuo*. The residue was subjected to flash chromatography (SiO₂, 9:1 CH₂Cl₂/2M NH₃ in MeOH) to afford
 15 0.49 g (77%) of the desired amine as a white foam. NMR (CDCl₃): 7.40 (dd, 1H), 7.37-7.20 (m, 6H), 7.07 (d, 1H), 6.95 (d, 1H), 6.15 (s, 2H), 5.66 (d, 1H), 5.06 (q, 1H), 4.06-3.63 (7H), 3.37-2.58 (m, 9H), 1.73-1.05 (m, 6H), 1.03 (s, 3H), 0.92 (s, 3H). MS (ESI): 634 (M+H).

20

Example (Compound 258)

Step 1:

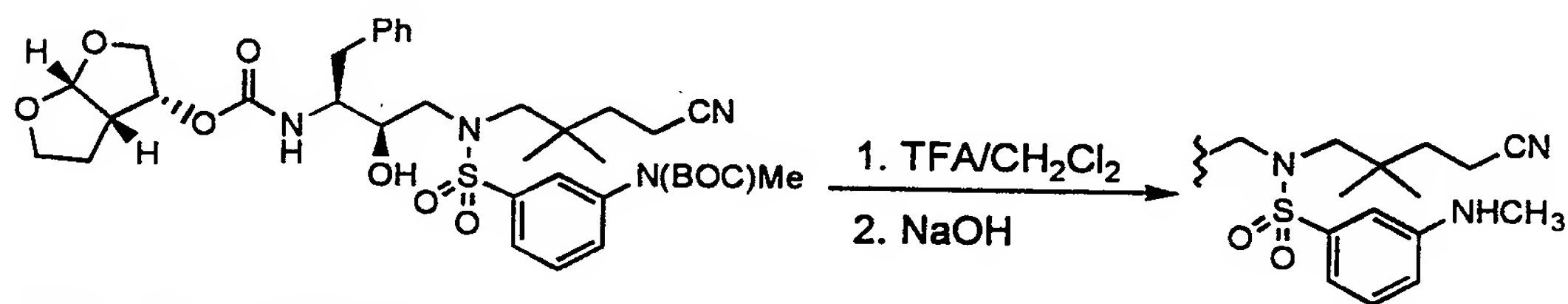


(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-1-benzyl-3-[(3-[(*tert*-butoxycarbonyl) (methyl) amino] phenylsulfonyl) (4-cyano-2,2-dimethylbutyl) amino]-2-
 25 hydroxypropylcarbamate

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A solution of 0.60 g (1.4 mmol) of (3*R*, 3*aS*, 6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-1-benzyl-3-[(4-cyano-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate (Example (Compound 257)), 0.83 g (2.7 mmol) of 3-[(*tert*-butoxycarbonyl)(methyl)amino]benzenesulfonyl chloride, 0.50 mL (2.8 mmol) of *N,N*-diisopropylethylamine, and 17 mg of DMAP in 15 mL of anhydrous THF was stirred at RT. After 18 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 97:3 to 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.36 g (36%) of the desired product as a light yellow foam. NMR (CDCl₃): 7.86 (s, 1H), 7.51 (m, 2H), 7.41 (d, 1H), 7.30-7.11 (m, 6H), 5.60 (d, 1H), 5.26 (m, 1H), 4.90 (q, 1H), 3.86 (m, 2H), 3.72 (m, 2H), 3.59 (m, 3H), 3.28 (s, 3H), 3.25 (d, 1H), 3.15 (m, 2H), 3.00 (d, 1H), 2.81 (m, 1H), 2.59 (dd, 1H), 2.39 (t, 2H), 1.78 (t, 2H), 1.54 (m, 1H), 1.52 (s, 9H), 1.26 (m, 1H), 1.00 (d, 6H). MS (ESI): 737 (M+Na).

Step 2:



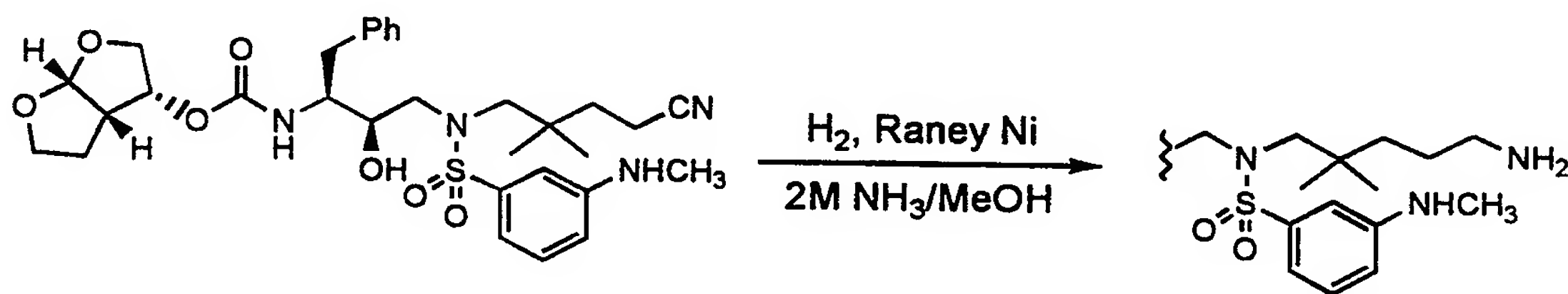
(3*R*, 3*aS*, 6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*, 2*R*)-1-benzyl-3-[(4-cyano-2,2-dimethylbutyl)[3-(methylamino)phenyl]sulfonylamino]-2-hydroxypropyl]carbamate

A solution of 0.35 g (0.49 mmol) of (3*R*, 3*aS*, 6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-1-benzyl-3-[(3-[(*tert*-butoxycarbonyl)(methyl)amino]phenylsulfonyl)(4-cyano-2,2-dimethylbutyl)amino]-2-

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hydroxypropylcarbamate in 20 mL of 1:1 TFA/CH₂Cl₂ was stirred at RT for 2 hours and then concentrated *in vacuo*. The residue was dissolved in CH₂Cl₂. The solution was washed with 1M aqueous NaOH (1x), aqueous brine (3x),
 5 dried over MgSO₄, and concentrated at reduced pressure to afford 0.29 g (95%) of the desired product as a colorless viscous oil. NMR (CDCl₃): 7.40–7.19 (m, 7H), 7.08 (d, 1H), 6.99 (s, 1H), 6.81 (dd, 1H), 5.68 (d, 1H), 5.02 (q, 1H), 4.88 (d, 1H), 4.23–3.62 (m, 7H), 3.34–2.84 (m, 8H), 2.79
 10 (dd, 1H), 2.42 (t, 2H), 1.80 (t, 2H), 1.61 (m, 1H), 1.42 (m, 1H), 1.01 (s, 6H). MS(ESI): 615 (M+H).

Step 3:

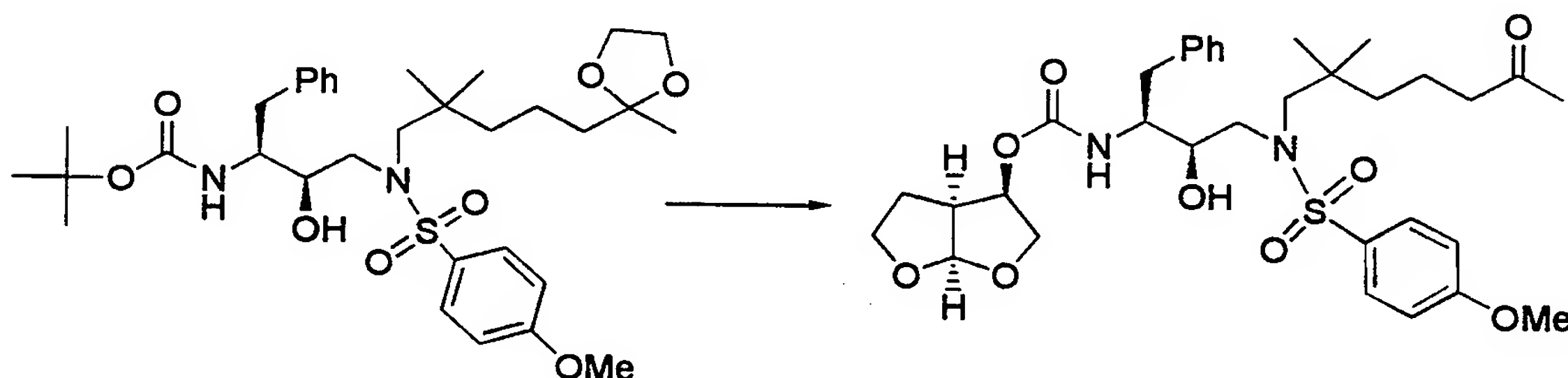


(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-
 ((5-amino-2,2-dimethylpentyl)[3-
 15 (methylamino)phenyl]sulfonylamino)-1-benzyl-2-
 hydroxypropyl]carbamate

A mixture of 0.29 g (0.46 mmol) of
 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-
 benzyl-3-((4-cyano-2,2-dimethylbutyl)[3-
 20 (methylamino)phenyl]sulfonylamino)-2-
 hydroxypropyl]carbamate and approximately 50 mg of Raney
 nickel (washed with water and MeOH) in 60 mL of 2M
 NH₃/MeOH was subjected to hydrogenation at 45 psi. After
 1.5 hours the reaction vessel was purged with nitrogen,
 25 catalyst removed by filtration through celite and the
 filtrate concentrated *in vacuo*. The residue was subjected

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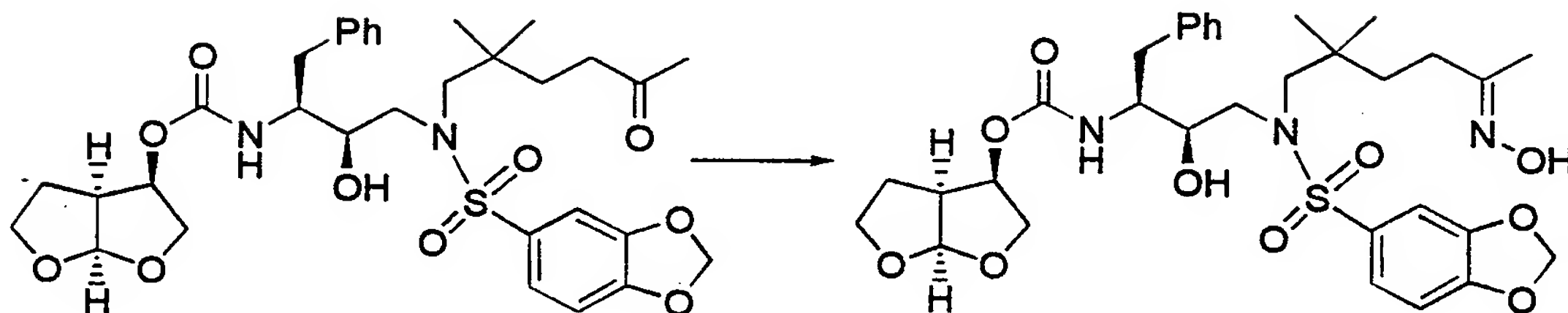
to flash chromatography (SiO₂, 9:1 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.22 g (76%) of the desired amine as a white foam. NMR (CDCl₃): 7.39-7.19 (m, 6H), 7.10 (d, 1H), 7.04-6.92 (m, 2H), 6.81 (dd, 1H), 5.63 (d, 1H), 5.05 (q, 1H), 4.16 (br s, 1H), 4.09-3.80 (m, 4H), 3.70 (m, 2H), 3.31 (m, 2H), 3.21-3.00 (m, 3H), 2.99-2.58 (m, 5H), 2.90 (s, 3H), 2.57-2.10 (br, 2H), 1.82-1.07 (m, 6H), 1.04 (s, 3H), 0.92 (s, 3H). MS(ESI): 619(M+H).

Example (Compound 259)

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-(2,2-dimethyl-5-oxoheptyl)[(4-methoxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

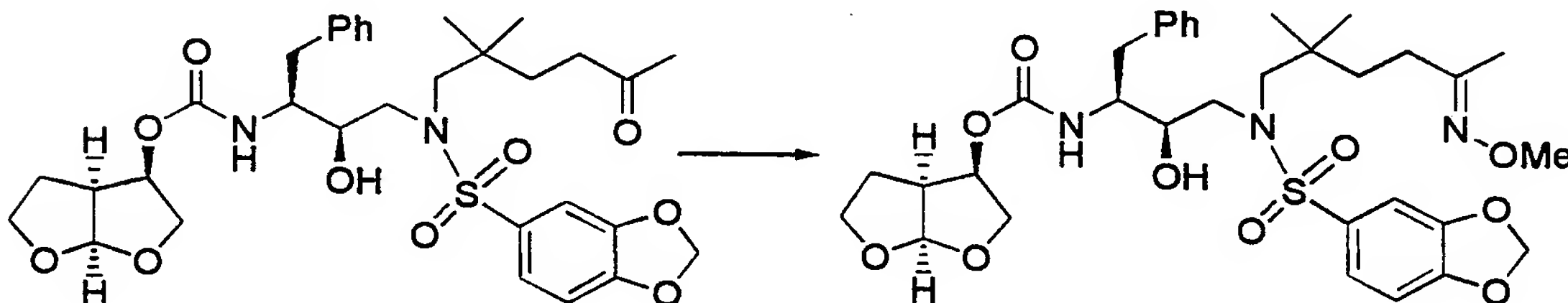
The product from step 3 (Example (Compound 209)) was subjected to procedures similar to those in Example (Compound 252) to give the title compound as a foam; ¹H NMR (DMSO-*d*₆): δ 0.88 (6H, s), 1.10-1.25 (4H, m), 1.3-1.5 (2H, m), 2.03 (3H, s), 2.25-2.35 (3H, m), 2.73 (1H, d), 2.79 (1H, t), 2.85-3.00 (2H, m), 3.2-3.3 (3H, m), 3.35-3.40 (1H, m), 3.55 (1H, dd), 3.6 (1H, dd), 3.65-3.75 (2H, m), 3.8 (3H, s), 4.8 (1H, dt), 5.05 (1H, d), 5.5 (1H, d), 7.1 (2H, d), 7.15-7.25 (6H, m), 7.75 (2H, d); MS: 669.1 (M+23); C₃₃H₄₆N₂O₉S.

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Example (Compound 260)

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-(2,2-dimethyl-5-hydroxyiminohexyl)[(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

The product from Example (Compound 240) was subjected to the procedure in Example (Compound 227) to afford the title compound as a foam; ^1H NMR (DMSO- d_6): δ 0.90 (6H, s), 1.1 (1H, dd), 1.20-1.45 (3H, m), 1.75 (3H, s), 2.05-2.10 (2H, m), 2.2 (1H, d), 2.7-2.8 (2H, m), 2.9-3.0 (2H, m), 3.25-3.35 (2H, m), 3.37-3.42 (1H, m), 3.55 (1H, d), 3.60 (1H, d), 3.7 (1H, t), 3.72-3.75 (1H, m), 3.79 (1H, d), 3.80 (1H, d), 4.8 (1H, dt), 5.1 (1H, d), 5.5 (1H, d), 6.2 (2H, s), 7.05 (1H, d), 7.10-7.25 (7H, m), 7.35 (1H, d); MS: 662.1 (MH^+); $\text{C}_{32}\text{H}_{43}\text{N}_3\text{O}_{10}\text{S}$.

Example (Compound 261)

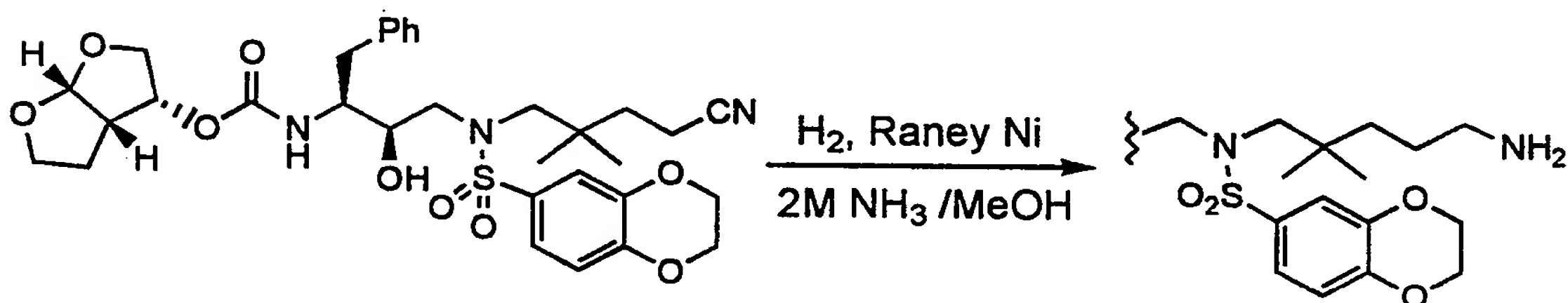
(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N-((1S,2R)-1-benzyl-3-(2,2-dimethyl-5-methoxyiminohexyl)[(3,4-

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methylenedioxyphenyl) sulfonyl] amino-2-hydroxypropyl) carbamate

The product from Example (Compound 240) was subjected to a procedure similar to that of Example (Compound 227), using methoxylamine hydrochloride to provide the title compound as a foam; ^1H NMR ($\text{DMSO}-d_6$): δ 0.90 (6H, s), 1.15 (1H, dd), 1.30-1.45 (3H, m), 1.75 (3H, s), 2.05-2.15 (2H, m), 2.2 (1H, d), 2.7-2.8 (2H, m), 2.9 (1H, dd), 2.95 (1H, d), 3.3-3.5 (3H, m), 3.52-3.60 (2H, m), 3.7 (3H, s), 3.75-3.83 (3H, m), 4.8 (1H, dt), 5.15 (1H, d), 5.5 (1H, d), 6.1 (2H, s), 7.05 (1H, d), 7.10-7.25 (7H, m), 7.35 (1H, d); MS: 676.2 (MH^+); $\text{C}_{33}\text{H}_{45}\text{N}_3\text{O}_{10}\text{S}$.

Example (Compound 262)



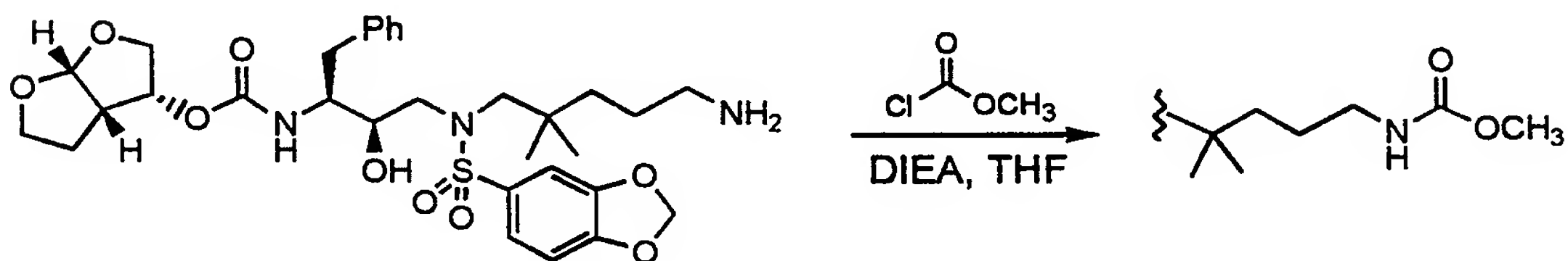
(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(5-amino-2,2-dimethylpentyl) (2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate

A mixture of 0.53 g (0.82 mmol) of (3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-cyano-2,2-dimethylbutyl) (2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) amino]-2-hydroxypropylcarbamate (obtained in a directly analogous way to that outlined in Example (Compound 257)) and approximately 50 mg of Raney nickel (washed with water and MeOH) in 60 mL of 2M NH_3/MeOH was subjected to hydrogenation at 45 psi. After

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2.0 hours the reaction vessel was purged with nitrogen, catalyst removed by filtration through celite and the filtrate concentrated *in vacuo*. The residue was subjected to flash chromatography (SiO₂, 9:1 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.46 g (87%) of the desired amine as a white foam. NMR (CDCl₃): 7.37-7.19 (m, 8H), 7.01 (m, 2H), 5.64 (d, 1H), 5.05 (q, 1H), 4.36 (m, 4H), 4.06-3.79 (m, 4H), 3.70 (m, 2H), 3.36-2.58 (m, 11H), 2.58-2.00 (br, 2H), 1.71-1.27 (m, 4H), 1.02 (s, 3H), 0.91 (s, 3H). MS (ESI): 648 (M+H).

Example (Compound 263)



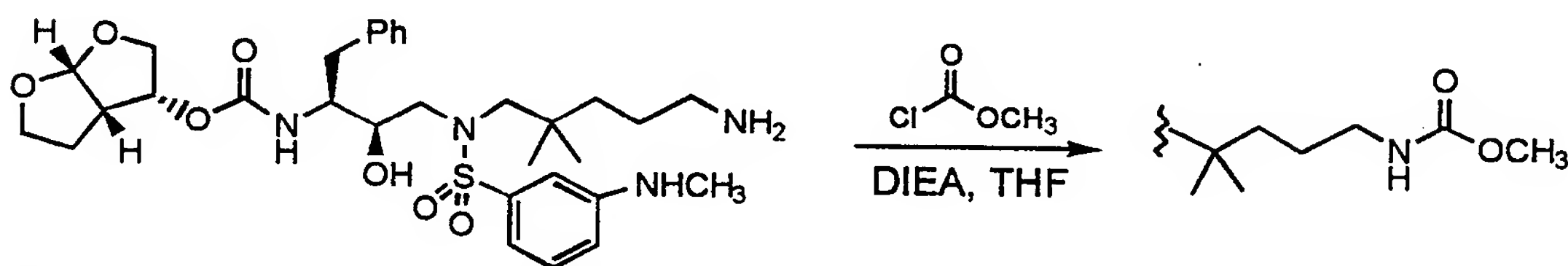
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-((1,3-benzodioxol-5-ylsulfonyl)amino)-2,2-dimethylpentylamino]-1-benzyl-2-hydroxypropyl carbamate

A solution of 60 mg (0.095 mmol) of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(5-amino-2,2-dimethylpentyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropyl carbamate and 20 µL (0.12 mmol) of N,N-diisopropylethylamine in 4 mL of anhydrous THF at 0°C was treated with 8.1 µL (0.11 mmol) of methyl chloroformate. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH)

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to afford 65 mg (98%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.40 (dd, 1H), 7.32-7.20 (m, 6H), 6.95 (d, 1H), 6.13 (s, 2H), 5.65 (d, 1H), 5.45 (d, 1H), 5.04 (m, 2H), 4.20-3.80 (m, 5H), 3.71 (m, 5H), 3.30-2.68 (m, 9H), 1.70-1.20 (m, 6H), 0.98 (s, 3H), 0.92 (s, 3H). MS (ESI): 714 (M+Na).

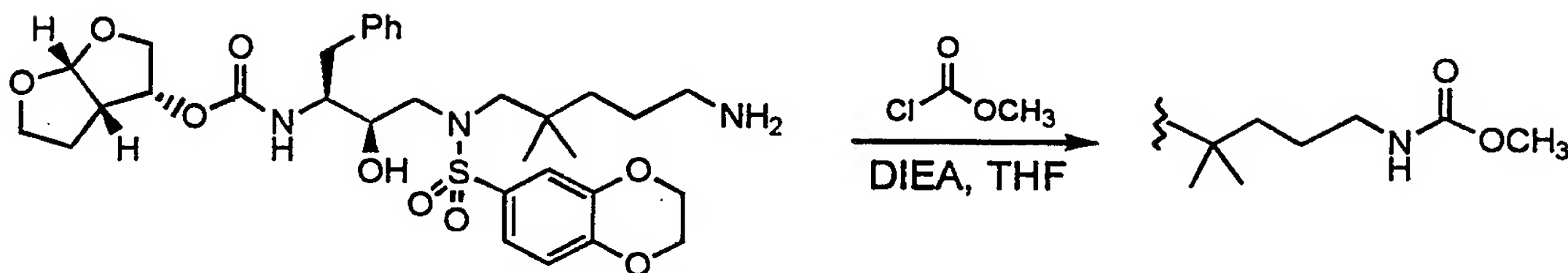
Example (Compound 264)



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-benzyl-2-hydroxy-3-(5-[(methoxycarbonyl)amino]-2,2-dimethylpentyl)[3-(methylanilino)phenyl]sulfonylamino)propyl]carbamate

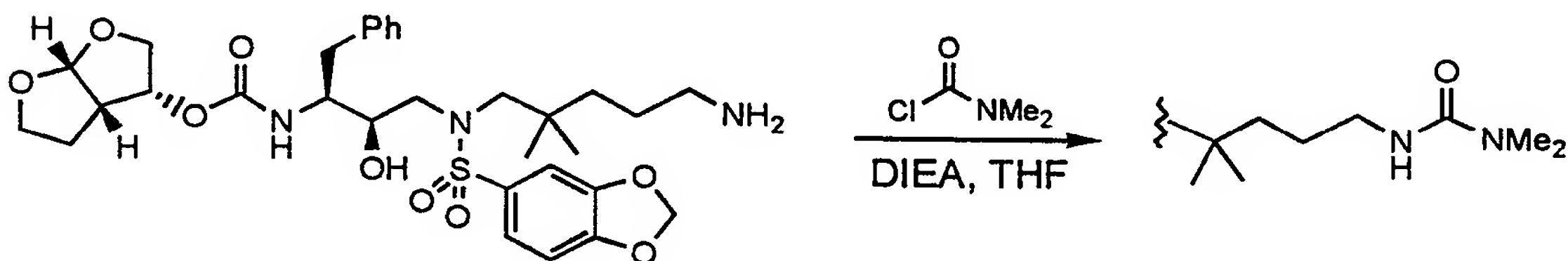
A solution of 50 mg (0.081 mmol) of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-((5-amino-2,2-dimethylpentyl)[3-(methylanilino)phenyl]sulfonylamino)-1-benzyl-2-hydroxypropyl]carbamate and 17 μ L (0.097 mmol) of N,N-diisopropylethylamine in 4 mL of anhydrous THF at 0°C was treated with 6.9 μ L (0.089 mmol) of methyl chloroformate. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated in vacuo and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 54 mg (98%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.28 (m, 6H), 6.99 (d, 1H), 6.89 (s, 1H), 6.71 (dd, 1H), 5.54 (d, 1H), 5.30 (d, 1H), 4.93 (m, 2H), 4.17-3.50 (m, 11H), 3.25-2.60 (m, 12H), 1.60-1.12 (m, 6H), 0.89 (s, 3H), 0.81 (s, 3H). MS (ESI): 677 (M+H).

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Example (Compound 265)

(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*, 2*R*)-1-benzyl-3-((2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl)5-[(methoxycarbonyl)amino]-2,2-dimethylpentylamino)-2-hydroxypropyl]carbamate

A solution of 60 mg (0.093 mmol) of (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-3-[(5-amino-2,2-dimethylpentyl) (2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate and 20 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 4 mL of anhydrous THF at 0°C was treated with 7.9 μ L (0.10 mmol) of methyl chloroformate. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 60 mg (91%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.38-7.19 (m, 7H), 7.01 (d, 1H), 5.64 (d, 1H), 5.42 (d, 1H), 5.02 (m, 2H), 4.36 (m, 4H), 4.19 (s, 1H), 4.12-3.80 (m, 4H), 3.70 (m, 5H), 3.30-2.70 (m, 9H), 1.61-1.22 (m, 6H), 0.98 (s, 3H), 0.91 (s, 3H). MS(ESI): 728 (M+Na).

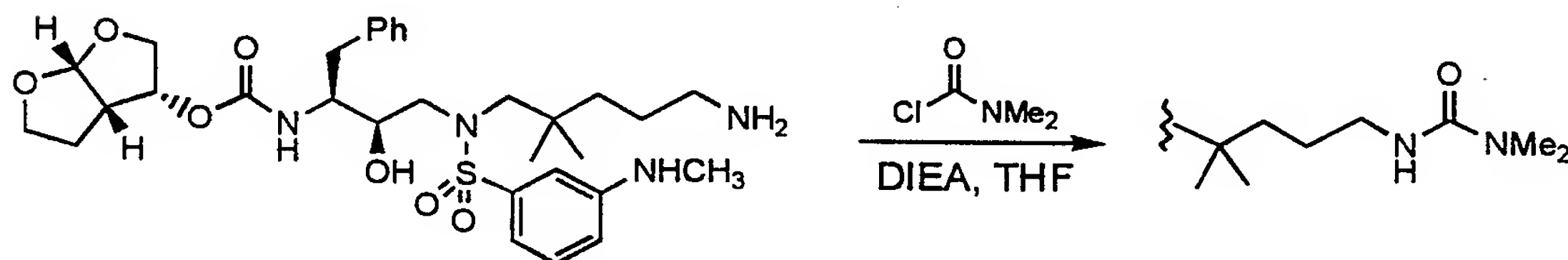
Example (Compound 266)

-252-

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl) (5-
 [(dimethylamino) carbonyl] amino-2,2-dimethylpentyl) amino]-
 1-benzyl-2-hydroxypropylcarbamate

5 A solution of 60 mg (0.095 mmol) of
 3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(5-amino-2,2-dimethylpentyl) (1,3-benzodioxol-5-
 ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate and
 20 μ L (0.11 mmol) of N,N-diisopropylethylamine in 5 mL of
 10 anhydrous THF at 0°C was treated with 9.6 μ L (0.11 mmol)
 of N,N-dimethylcarbamy l chloride. The solution was
 allowed to warm to RT with stirring. After 18 hours the
 solution was concentrated in vacuo and the residue was
 subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃
 15 in MeOH) to afford 55 mg (82%) of the desired compound as
 a white foam. H¹-NMR (CDCl₃): 7.35 (d, 1H), 7.27-7.11 (m,
 6H), 6.88 (d, 1H), 6.08 (s, 2H), 6.03 (d, 1H), 5.60 (d,
 1H), 4.94 (q, 1H), 4.63 (br s, 1H), 3.89 (m, 2H), 3.80
 (m, 2H), 3.62 (m, 2H), 3.32-3.01 (m, 6H), 2.88 (s, 6H),
 20 2.81 (m, 3H), 2.69 (dd, 1H), 1.58-1.17 (m, 6H), 0.95 (s,
 3H), 0.86 (s, 3H). MS (ESI): 705 (M+H).

Example (Compound 267)



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-1-
 benzyl-3-((5-[(dimethylamino) carbonyl] amino-2,2-

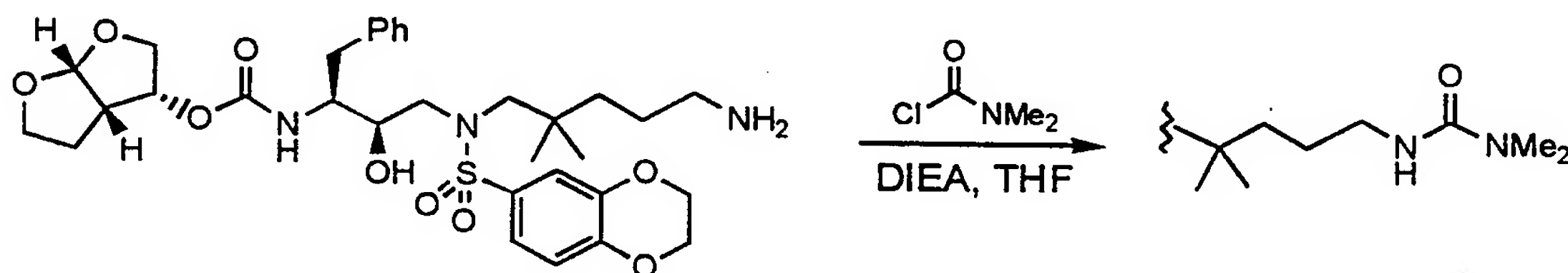
-253-

dimethylpentyl) [3- (methylamino)phenyl]sulfonylamino) -2-hydroxypropyl] carbamate

A solution of 50 mg (0.081 mmol) of (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*, 2*R*)-3-((5-amino-2,2-dimethylpentyl) [3- (methylamino)phenyl]sulfonylamino)-1-benzyl-2-hydroxypropyl]carbamate and 17 μ L (0.097 mmol) of *N,N*-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 8.2 μ L (0.089 mmol) of *N,N*-dimethylcarbamylyl chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 44 mg (79%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.31-7.11 (m, 6H), 7.03 (d, 1H), 6.95 (s, 1H), 6.73 (d, 1H), 5.92 (d, 1H), 5.60 (d, 1H), 4.94 (q, 1H), 4.63 (t, 1H), 4.19 (m, 2H), 3.94-3.73 (m, 4H), 3.63 (m, 2H), 3.30-3.00 (m, 6H), 2.91-2.78 (m, 11H), 2.69 (dd, 1H), 1.59-1.19 (m, 6H), 0.94 (s, 3H), 0.86 (s, 3H). MS(ESI): 690 (M+H).

20

Example (Compound 268)

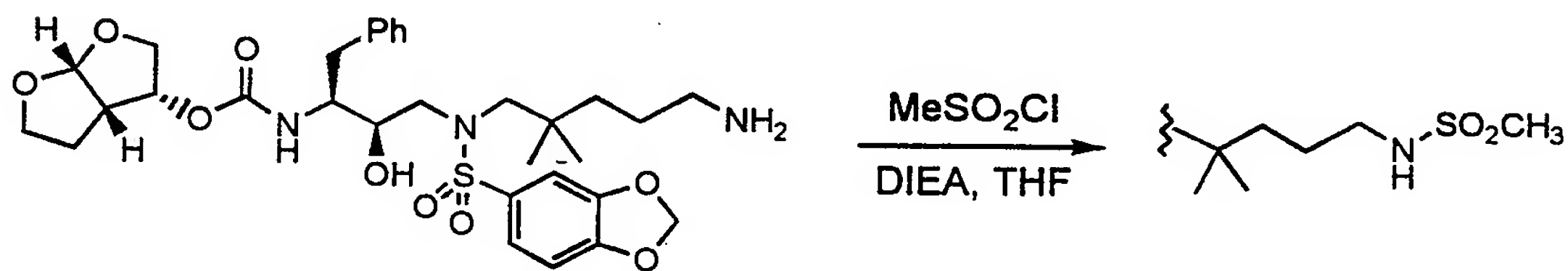


(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-1-benzyl-3-[(2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) (5-[(dimethylamino) carbonyl]amino-2,2-dimethylpentyl) amino]-2-hydroxypropylcarbamate

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A solution of 60 mg (0.093 mmol) of (3*R*, 3*aS*, 6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-3-[(5-amino-2,2-dimethylpentyl) (2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate and 20 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 9.4 μ L (0.10 mmol) of *N,N*-dimethylcarbamy chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated in vacuo and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 54 mg (81%) of the desired compound as a white foam. H¹-NMR (CDCl₃): 7.31-7.12 (m, 7H), 6.96 (d, 1H), 6.02 (d, 1H), 5.60 (d, 1H), 4.95 (q, 1H), 4.62 (t, 1H), 4.34-4.21 (m, 5H), 3.91 (m, 2H), 3.80 (m, 2H), 3.62 (m, 2H), 3.32-2.99 (m, 6H), 2.89 (s, 6H), 2.80 (m, 2H), 2.69 (dd, 1H), 1.58-1.18 (m, 6H), 0.93 (s, 3H), 0.86 (s, 3H). MS(ESI): 719 (M+H).

Example (Compound 269)



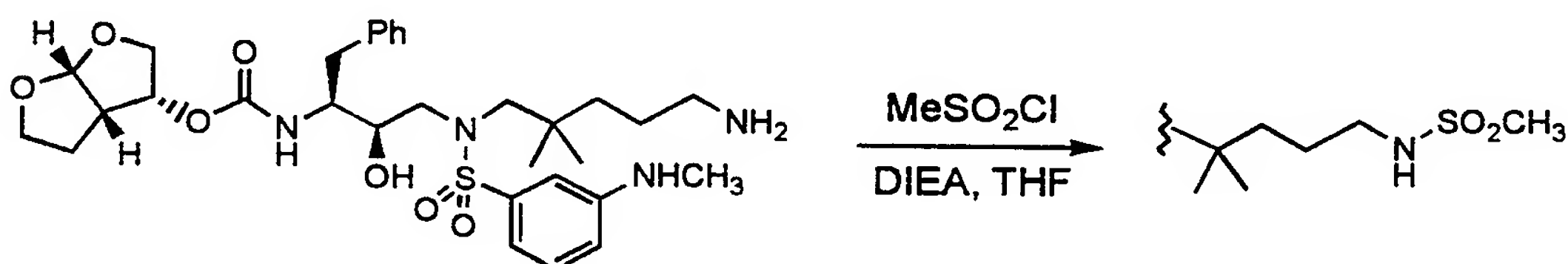
(3*R*, 3*aS*, 6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*, 2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) 2,2-dimethyl-5-[(methylsulfonyl) amino]pentylamino)-1-benzyl-2-hydroxypropyl]carbamate

A solution of 60 mg (0.095 mmol) of 3*R*, 3*aS*, 6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*, 2*R*)-3-[(5-amino-2,2-dimethylpentyl) (1,3-benzodioxol-5-

-255-

ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate and 20 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 8.1 μ L (0.11 mmol) of methanesulfonyl chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated *in vacuo* and the residue was subjected to flash chromatography (SiO_2 , 95:5 CH_2Cl_2 /2M NH_3 in MeOH) to afford 66 mg (97%) of the desired compound as a white foam. $^1\text{H-NMR}$ (CDCl_3): 7.33 (d, 1H), 7.29-7.14 (m, 6H), 6.89 (d, 1H), 6.09 (s, 2H), 5.60 (d, 1H), 5.14 (d, 1H), 4.96 (m, 2H), 4.04 (m, 2H), 3.90 (dd, 1H), 3.81 (m, 2H), 3.68 (m, 2H), 3.21-3.02 (m, 5H), 2.96 (d, 1H), 2.91 (s, 3H), 2.89-2.66 (m, 3H), 1.55 (m, 3H), 1.47-1.28 (m, 3H), 0.95 (s, 3H), 0.90 (s, 3H). MS (ESI): 734 (M+Na).

15

Example (Compound 270)

(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*,2*R*)-1-benzyl-3-((5-[(methanesulfonyl)amino]-2,2-dimethylpentyl)[3-(methylamino)phenyl]sulfonylamino)-2-hydroxypropyl]carbamate

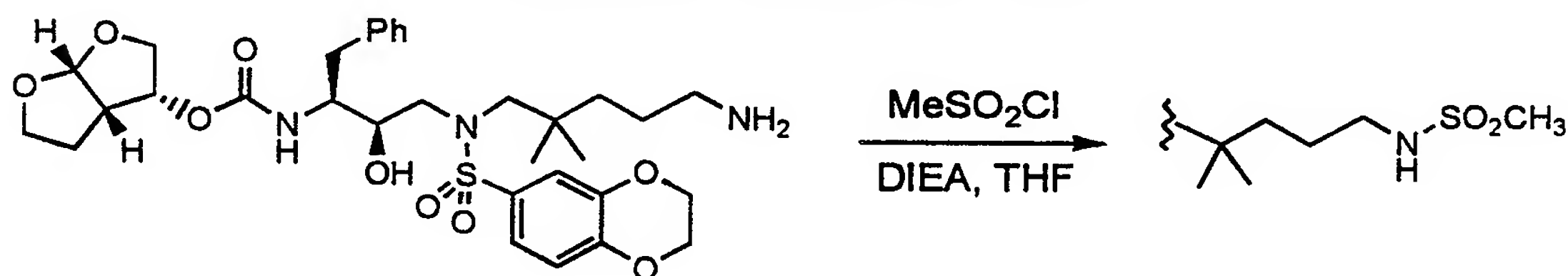
20

A solution of 50 mg (0.081 mmol) of (3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*,2*R*)-3-((5-amino-2,2-dimethylpentyl)[3-(methylamino)phenyl]sulfonylamino)-1-benzyl-2-hydroxypropyl]carbamate and 17 μ L (0.097 mmol) of *N,N*-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 6.9 μ L (0.089 mmol) of methanesulfonyl

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chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated in vacuo and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 55 mg (98%) of the desired compound as a white foam. H¹-NMR (CDCl₃): 7.32-7.12 (m, 6H), 7.01 (d, 1H), 6.92 (s, 1H), 6.78 (dd, 1H), 5.60 (d, 1H), 5.11 (d, 1H), 5.01-4.88 (m, 2H), 4.12 (br s, 1H), 4.04 (m, 2H), 3.93-3.77 (m, 3H), 3.65 (m, 2H), 3.25 (dd, 1H), 3.20-2.90 (m, 5H), 2.91 (s, 3H), 2.90-2.81 (m, 5H), 2.71 (dd, 1H), 1.58 (m, 3H), 1.49-1.28 (m, 3H), 0.96 (s, 3H), 0.89 (s, 3H). MS(ESI): 719 (M+Na).

Example (Compound 271)

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-((2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl)2,2-dimethyl-5-[(methanesulfonyl)amino]pentylamino)-1-benzyl-2-hydroxypropyl]carbamate

A solution of 60 mg (0.093 mmol) of (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(5-amino-2,2-dimethylpentyl)(2,3-dihydro-1,4-benzodioxin-6-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate and 20 µL (0.11 mmol) of N,N-diisopropylethylamine in 5 mL of anhydrous THF at 0°C was treated with 7.9 µL (0.10 mmol) of methanesulfonyl chloride. The solution was allowed to warm to RT with stirring. After 18 hours the solution was concentrated

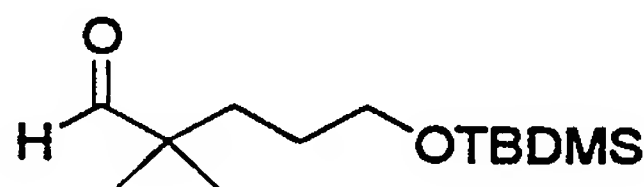
-257-

in vacuo and the residue was subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 64 mg (94%) of the desired compound as a white foam. ¹H-NMR (CDCl₃): 7.30-7.13 (m, 7H), 6.97 (d, 1H), 5.60 (d, 1H), 5.12 (d, 1H), 5.01-4.89 (m, 2H), 4.32 (m, 4H), 4.10 (s, 1H), 4.04 (m, 1H), 3.89 (dd, 1H), 3.81 (m, 2H), 3.65 (m, 2H), 3.21-3.02 (m, 5H), 2.95 (d, 1H), 2.92 (s, 3H), 2.88-2.65 (m, 3H), 1.55 (m, 3H), 1.45-1.27 (m, 3H), 0.94 (s, 3H), 0.89 (s, 3H). MS (ESI): 748 (M+Na).

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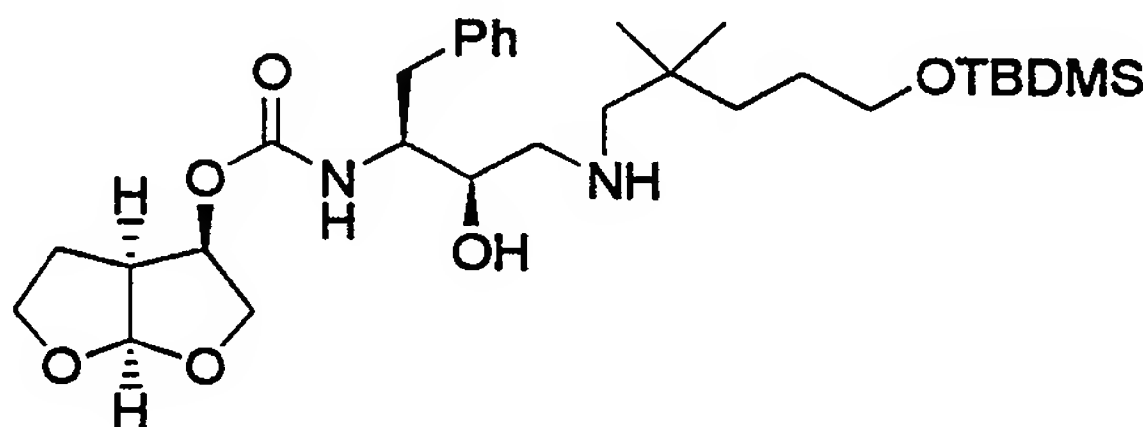
Example (Compound 272)

Step 1:

**5-tert-Butyldimethylsilyloxy-2,2-dimethylpentanal**

The title compound was prepared by procedure b (Example (Compound 24)) using 3-tert-butyltrimethylsilyloxybromopropane as alkylating agent. ¹H NMR (CDCl₃): δ -0.02 (6H, s), 0.84 (9H, s), 1.04 (6H, s), 1.38-1.42 (2H, m), 1.46-1.50 (2H, m), 3.55 (2H, t), 9.43 (1H, s); C₁₃H₂₈O₂Si.

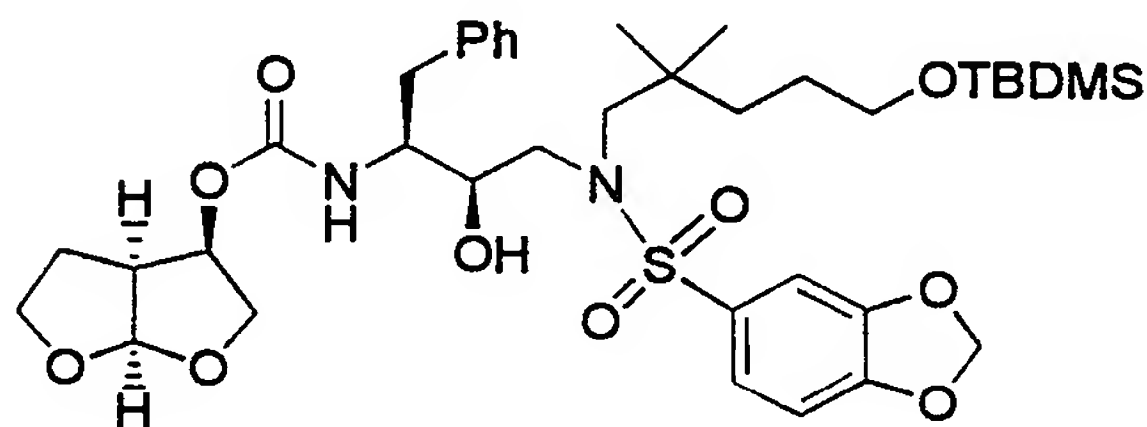
Step 2:



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(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-[(5-tert-butyldimethylsilyloxy-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate

To a suspension of (3R,3aS,6aR)-
5 hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-amino-1-benzyl-2-hydroxypropylcarbamate (0.5 g, 1.5 mmol) in N,N-dimethylformamide (10 mL) and 1,2-dichloroethane (10 mL) was added a solution of the product from step a (0.33 g, 1.3 mmol) in tetrahydrofuran (10 mL). Acetic acid (0.3
10 mL) was added followed by sodium triacetoxyborohydride (0.32 g, 1.5 mmol) and the resulting mixture was stirred at ambient temperature under nitrogen atmosphere for 16 hours. The mixture was concentrated and 0.5 N sodium hydroxide/water (40 mL) was added. The mixture was
15 extracted with ethyl acetate and the organic phase was washed with brine, dried (magnesium sulfate), and concentrated. The residue was chromatographed (silica gel, chloroform/methanol/ammonium hydroxide, 95:5:1) to provide the title compound as a white solid (0.61 g,
20 89%); ¹H NMR (DMSO-d₆): δ -0.02 (6H, s), 0.80 (6H, s), 0.83 (9H, s), 1.15-1.20 (2H, m), 1.25-1.50 (4H, m), 2.25 (2H, dd), 2.45-2.52 (3H, m), 2.57 (1H, d), 2.76 (1H, quartet), 3.0 (1H, dd), 3.45 (1H, br s), 3.5-3.6 (5H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.8-4.9 (2H, m), 5.5 (1H, d),
25 7.10-7.25 (6H, m); MS: 565.3 (MH⁺); C₃₀H₅₂N₂O₆Si.



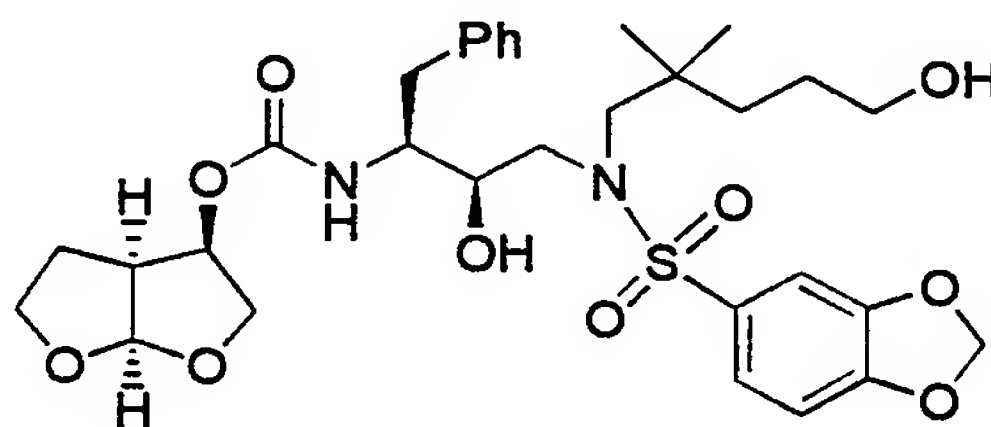
Step 3:

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(3*R*, 3*aS*, 6*aR*) -Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-1-benzyl-3-(5-*tert*-butyldimethylsilyloxy-2,2-dimethylpentyl)[(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

5 To a solution of the product from step b (0.2 g, 0.35 mmol) in dichloromethane (3 mL) at 0°C was added *N*-ethyldiisopropylamine (0.12 mL, 0.71 mmol) and 3,4-methylenedioxyphenylsulfonyl chloride (0.094 g, 0.42 mmol). The mixture was stirred at ambient temperature for
 10 72 hours, concentrated and chromatographed (silica gel, hexane/ethyl acetate, 3:2) to afford the title compound (0.22 g, 83%) as a white foam; ¹H NMR (DMSO-*d*₆): δ -0.02 (6H, s), 0.82 (9H, s), 0.88 (6H, s), 1.15 (1H, dd), 1.20-1.43 (4H, m), 2.4 (1H, dd), 2.5-2.6 (2H, m), 2.9-3.0 (2H, m), 3.30-3.38 (3H, m), 3.39-3.42 (1H, m), 3.5-3.6 (4H, m), 3.67 (1H, t), 3.75 (1H, quartet), 3.8 (1H, dd), 4.8 (1H, dt), 5.1 (1H, d), 5.45 (1H, d), 6.15 (2H, s), 7.02 (1H, d), 7.1-7.3 (7H, m), 7.34 (1H, d); C₃₇H₅₆N₂O₁₀SSi.

Step 4:

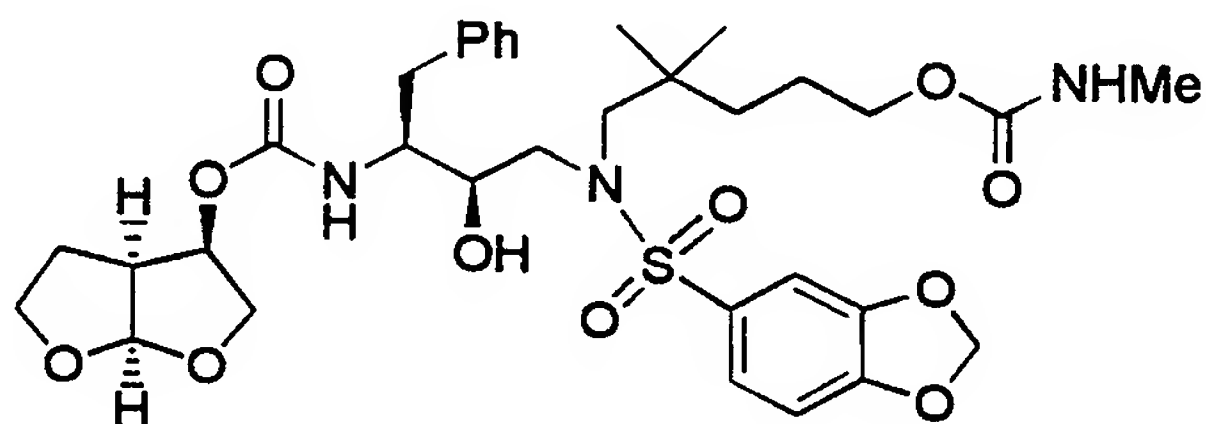


20 (3*R*, 3*aS*, 6*aR*) -Hexahydrofuro[2,3-*b*]furan-3-yl *N*-((1*S*, 2*R*)-1-benzyl-3-(2,2-dimethyl-5-hydroxypentyl)[(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

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A solution of the product from step c (0.19 g, 0.26 mmol) in tetrahydrofuran (2 mL), acetic acid (2 mL), and water (0.65 mL) was stirred at ambient temperature for 16 hours. The mixture was poured into saturated sodium bicarbonate/water and extracted with diethyl ether (3x). The combined organic layers were washed with water, dried (magnesium sulfate), concentrated, and dried *in vacuo* to provide the title compound (~0.19 g) which was used in the next step without further purification.

Step 6:



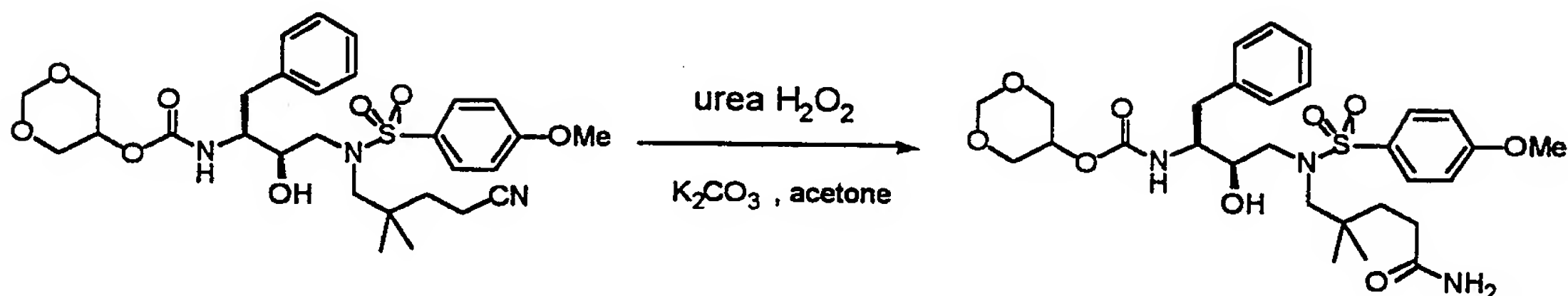
(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-benzyl-3-[2,2-dimethyl-5-(N'-methylcarbamoyloxypropyl)][(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl) carbamate

A solution of the product from step d (~0.19 g) and methyl isocyanate (0.36 mL, 6.1 mmol) in dichloromethane (2 mL) was stirred at ambient temperature for 64 hours. Solvent was evaporated and the residue was chromatographed (silica gel, chloroform/methanol, 98:2) to provide the title compound (0.11 g, 64%) as a white foam; ¹H NMR (DMSO-d₆): δ 0.9 (3H, s), 0.92 (3H, s), 1.13 (1H, dd), 1.20-1.35 (2H, m), 1.45-1.55 (2H, m), 2.2 (1H, dd), 2.54 (3H, d), 2.7-2.8 (2H, m), 2.9 (1H, dd), 2.95 (1H, dd), 3.30-3.43 (5H, m), 3.5-3.6 (2H, m), 3.7 (1H, td), 3.75 (1H, br quartet), 3.85 (1H, dd), 3.88 (1H, t), 4.8 (1H, dt), 5.1 (1H, d), 5.5 (1H, d), 6.15 (2H, s), 6.9

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(1H, quartet), 7.03 (1H, d), 7.1-7.25 (7H, m), 7.32 (1H, d); MS: 692.1 (MH⁺); C₃₃H₄₅N₃O₁₁S.

Example (Compound 273)

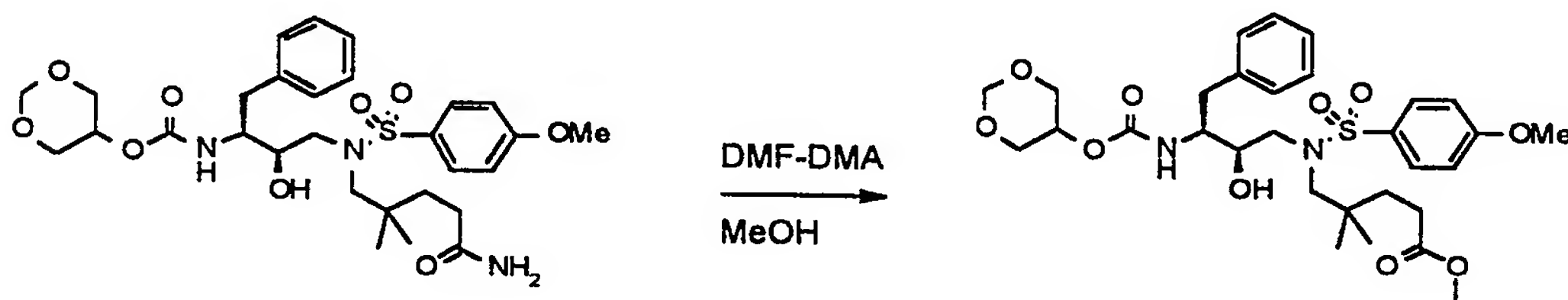


5 { (1S,2R)-1-Benzyl-3-[(4-carboxamide-2,2-dimethyl-butyl)-(4-methoxy-benzenesulfonyl)-amino]-2-hydroxy-propyl}-carbamic acid [1,3] dioxan-5-yl ester.

{ (1S,2R)-1-Benzyl-3-[(4-cyano-2,2-dimethyl-butyl)-(4-methoxy-benzenesulfonyl)-amino]-2-hydroxy-propyl}-carbamic acid [1,3] dioxan-5-yl ester (0.042 g, 0.071 mmol) was dissolved in 2 mL of 1:1, acetone : water and treated with urea hydroperoxide (0.20 mg, 0.212 mmol) at ambient temperature under argon with stirring. After 15h the reaction mixture was concentrated *in vacuo*, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed *in vacuo*. Purification by preparatory tlc (5% MeOH in CH₂Cl₂, SiO₂) gave 0.014 g of a white solid. MS (ES): 608 (M+1).

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Example (Compound 274)



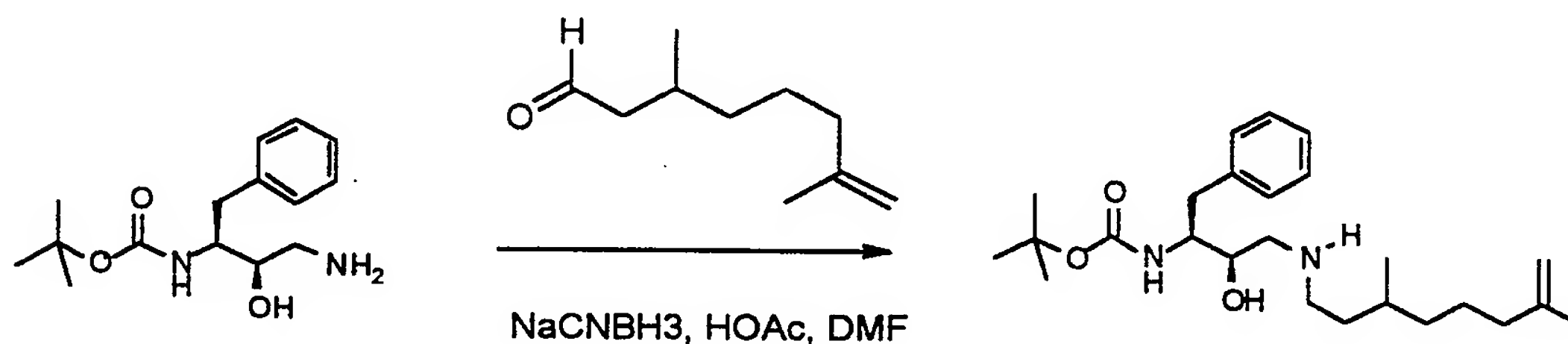
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{ (1S,2R)-1-Benzyl-3-[(4-carbomethoxy-2,2-dimethyl-butyl)-(4-methoxy-benzenesulfonyl)-amino]-2-hydroxy-propyl}-carbamic acid [1,3] dioxan-5-yl ester.

{ (1S,2R)-1-Benzyl-3-[(4-carboxamide-2,2-dimethyl-butyl)-(4-methoxy-benzenesulfonyl)-amino]-2-hydroxy-propyl}-carbamic acid [1,3] dioxan-5-yl ester (0.0063 g, 0.01 mmol) was dissolved in 1 mL of methanol and treated with dimethyl formamide-dimethyl acetal (0.20 mL, 3 EQ.) at ambient temperature under argon with stirring. After 15h the reaction mixture was concentrated in vacuo, taken up in EtOAc, washed with sat. aq. NaHCO₃, and brine. The organic phase was dried over MgSO₄, filtered and solvent removed in vacuo. Purification by preparatory tlc (5% MeOH in CH₂Cl₂) provided 2.5 mg of the derived methyl ester, MS(ES)= 623 amu (M+1).

Example (Compound 275)

Step 1:



20 [(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3, 7-dimethyloct-7-nyl)butyl amine].

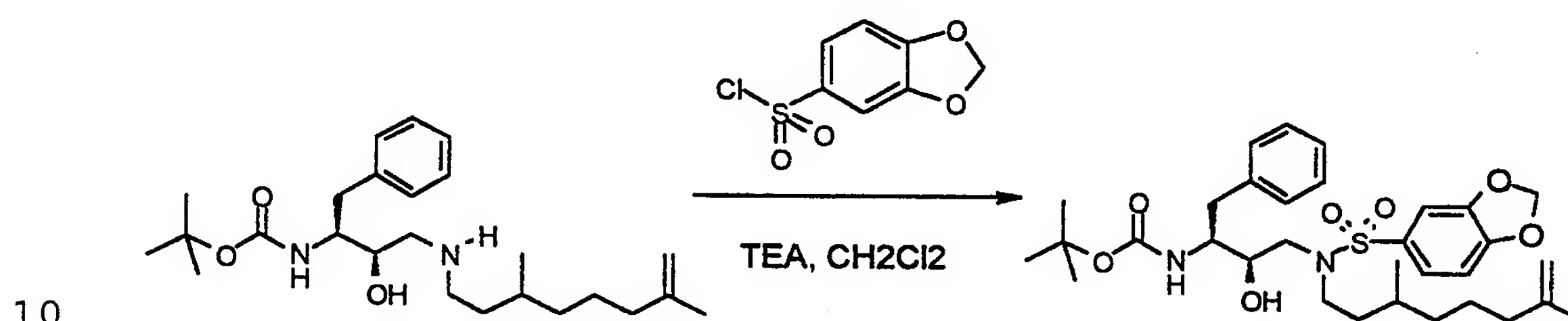
[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino) butane] (0.94 g, 3.35 mmol) was added to a solution of racemic 3, 7-dimethyloct-7-nyl carboxaldehyde in DMF (5 mL). This was followed by the addition of approximately one mL of glacial acetic acid and then NaCNBH₃ (2.0g, 9.6 mmol). The reaction was

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stirred under N₂ for 15 hrs, followed by diluting with 100 mL of CH₂Cl₂. The reaction mixture was then washed with sat. NaHCO₃, and brine. The organic phase was dried with MgSO₄ and the solvent was removed *in vacuo*.

- 5 Purification by SiO₂ chromatography (5% MeOH/CH₂Cl₂). Isolated 0.055 g of the product as a colorless residue: R_f=0.50 (5% MeOH/CH₂Cl₂).

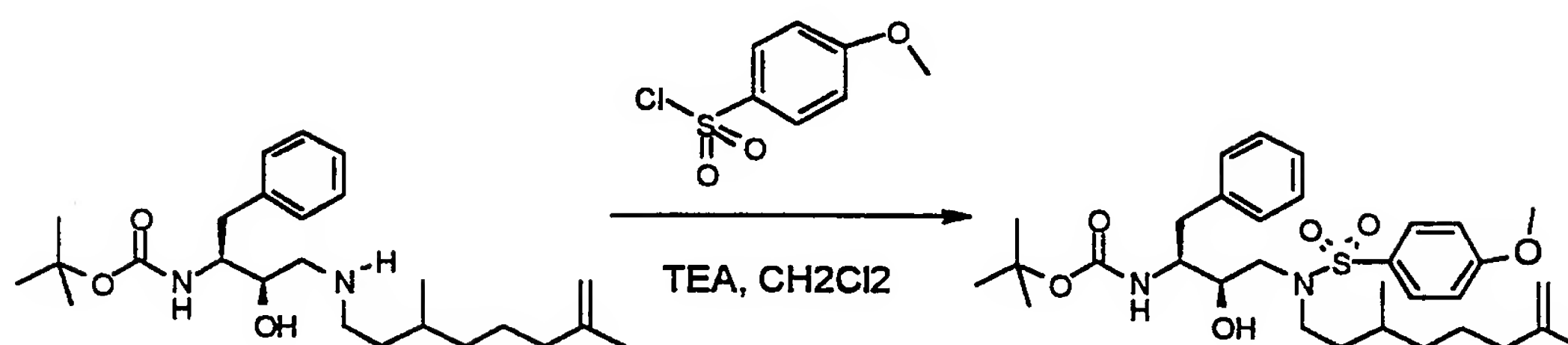
Step 2:



[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3,7-dimethyloct-7-enyl)] [2-[(Benzo[1,3]dioxole-5-sulfonyl)]butylamine.

- 15 [(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3, 7-dimethyloct-7-enyl)butyl amine] (27.6 mg, 0.066 mmol) was dissolved in 1 mL of CH₂Cl₂ followed by [2-[(Benzo[1,3]dioxole-5-sulfonyl chloride (15 mg, 0.066 mmol). TEA (200 uL) was then added and the reaction was stirred for 14 hours at room temperature. The reaction
- 20 was quenched by the addition of 3 mL of saturated NaHCO₃, followed by washing with sat. NaHCO₃, and then brine. The organic phase was dried with MgSO₄ and the solvent was removed *in vacuo*. Purification by SiO₂ chromatography (5% MeOH/ CH₂Cl₂). Recovered 0.0038 g of
- 25 the product as a colorless residue: R_f=0.50 (5% MeOH/ CH₂Cl₂/SiO₂), MS(ES) = 647 amu (M+1).

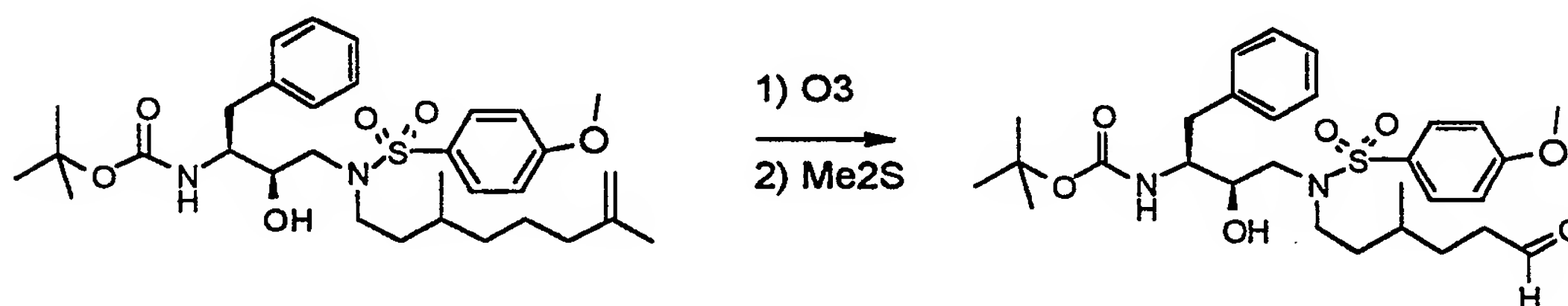
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Example (Compound 276)

[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3,7-dimethyloct-7-enyl)] [4-methoxyphenylsulfonyl]

5 butylamine.

[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3,7-dimethyloct-7-enyl)butyl amine] (27.6 mg, 0.066 mmol) was dissolved in 1 mL of CH₂Cl₂ followed by the addition of 4-methoxyphenylsulfonyl chloride (13.7 mg, 0.066 mmol). TEA (200 μ L) was then added and the reaction was stirred for 14 hours at room temperature. The reaction was quenched by the addition of 3 mL of saturated NaHCO₃, followed by washing with sat. NaHCO₃, and then brine. The organic phase was dried with MgSO₄ and the solvent was removed *in vacuo*. Purification by SiO₂ chromatography (5% MeOH/ CH₂Cl₂). Recovered 0.0021 g of the product as a colorless residue: R_f=0.50 (5% MeOH/CH₂Cl₂/SiO₂), MS(ES) = 633 amu (M+1).

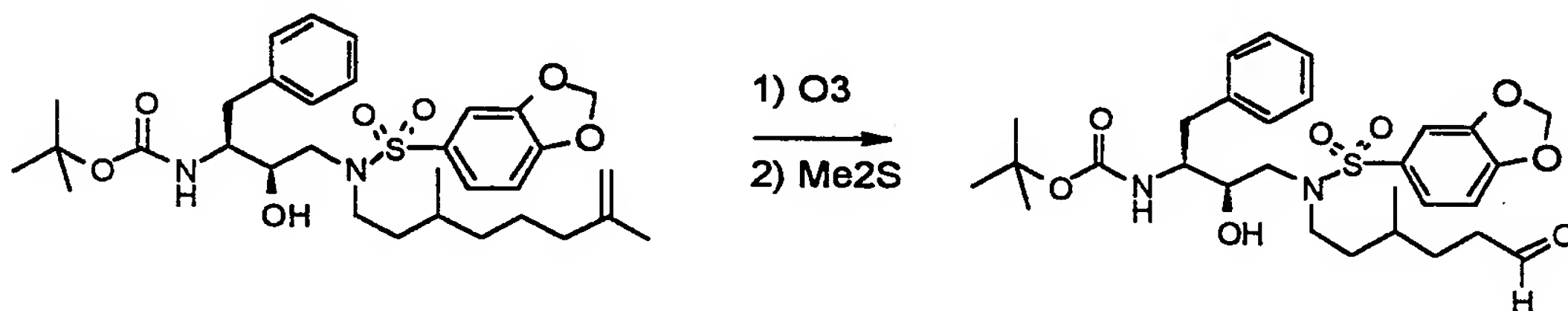
Example (Compound 277)

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[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3,7-dimethylheptylcarboxaldehyde)[4-methoxyphenylsulfonyl]butylamine.

[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3,7-dimethyloct-7-enyl)][4-methoxyphenylsulfonyl]butylamine (0.9 mg) was dissolved in 1 mL of CH₂Cl₂ containing 100 μ L of MeOH. The reaction mixture was cooled to -78 °C under a stream of O₂. A stream of ozone was then bubbled through the reaction mixture for five minutes, until a pale blue color was evident. The reaction was then purged of ozone by bubbling O₂ through for five minutes and then N₂ for an additional 5 minutes and the addition of 200 μ L of dimethyl sulfide. After stirring for 30 minutes at -78 °C the reaction was warmed to room temperature and concentrated *in vacuo*. The reaction mixture was purified by prep. TLC (SiO₂) eluting with 5% MeOH/CH₂Cl₂, 0.5 mg of the desired aldehyde was isolated; MS(ES) = 563 amu (M+1). Note that olefin migration occurred during the reaction conditions prior to exposure to .

Example (Compound 278)



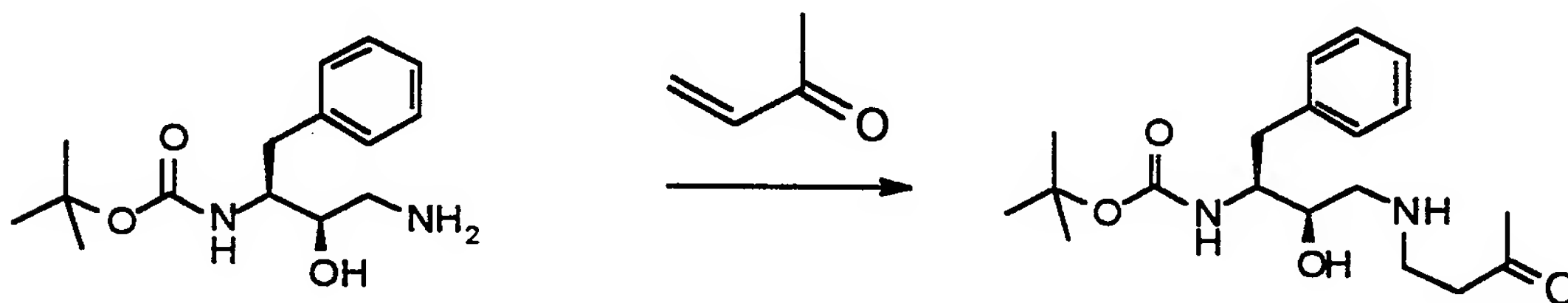
[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3,7-dimethylheptylcarboxaldehyde[(Benzo[1,3]dioxole-5-sulfonyl]butylamine.

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[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(3,7-dimethyloct-7-enyl)] [2-
 [(Benzo[1,3]dioxole-5-sulfonyl]butylamine (4.0 mg) was
 dissolved in 1 mL of CH₂Cl₂ containing 100 uL of MeOH and
 5 1 mg K₂CO₃. The reaction mixture was cooled to -78 °C
 under a stream of O₂. A stream of ozone was then bubbled
 through the reaction mixture for five minutes, until a
 pale blue color was evident. The reaction was then
 purged of ozone by bubbling O₂ through for five minutes,
 10 followed by the addition of 200 uL of dimethyl sulfide.
 After stirring for 30 minutes at -78 °C the reaction was
 warmed to room temperature and concentrated *in vacuo*.
 The reaction mixture was purified by prep. TLC (SiO₂)
 eluting with 5% MeOH/CH₂Cl₂, 2.1 mg of the desired
 15 aldehyde was isolated; MS(ES) = 577 amu (M+1), 621 amu
 for (M-1+ TFA). Note that olefin migration occurred
 during the reaction conditions prior to exposure to ozone
 both with and without base present.

Example (Compound 279)

20 Step 1:



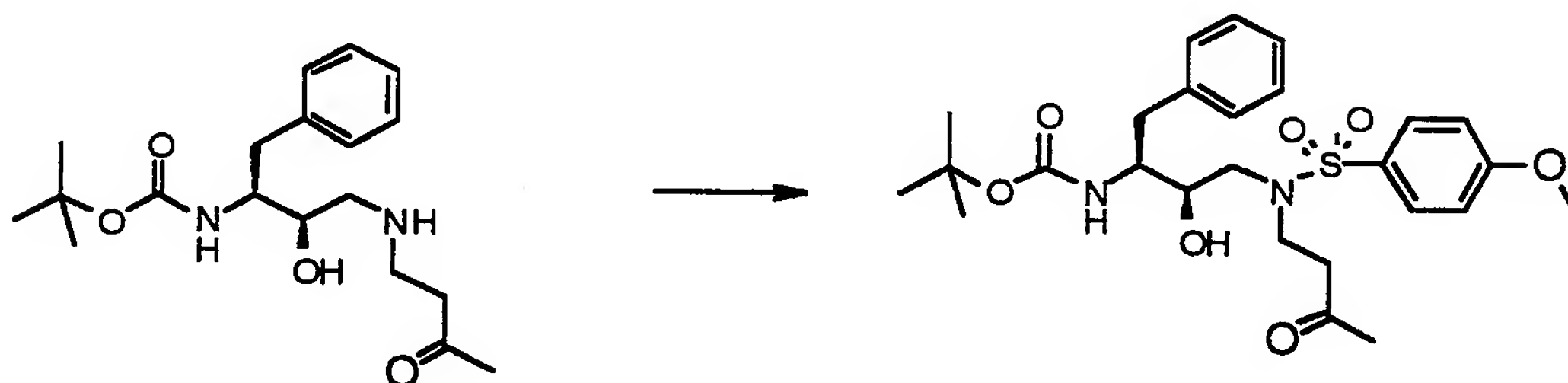
[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-
 (1-amino-3-butanone) butane].

[(2R,3S)-3-*tert*-Butoxycarbonylamino-2-hydroxy-
 25 4-phenyl-(1- amino) butane] (0.268 g, 1.0 mmol) was
 dissolved in EtOH (5 mL) and cooled to 0°C. This was
 followed by the addition methyl vinyl ketone (79.9 uL,

-267-

1.0 mmol). The reaction was stirred under N₂ for 15 hrs and gradually warmed to room temperature as the ice bath melted. The reaction mixture was then concentrated in vacuo and purified by silica gel chromatography (gradient from 2 % MeOH/ CH₂Cl₂ to 90% MeOH/CH₂Cl₂ to 85/10/5 MeOH/CH₂Cl₂/NH₄OH. Isolated both unreacted starting material, the mono (desired) and the bis addition product.

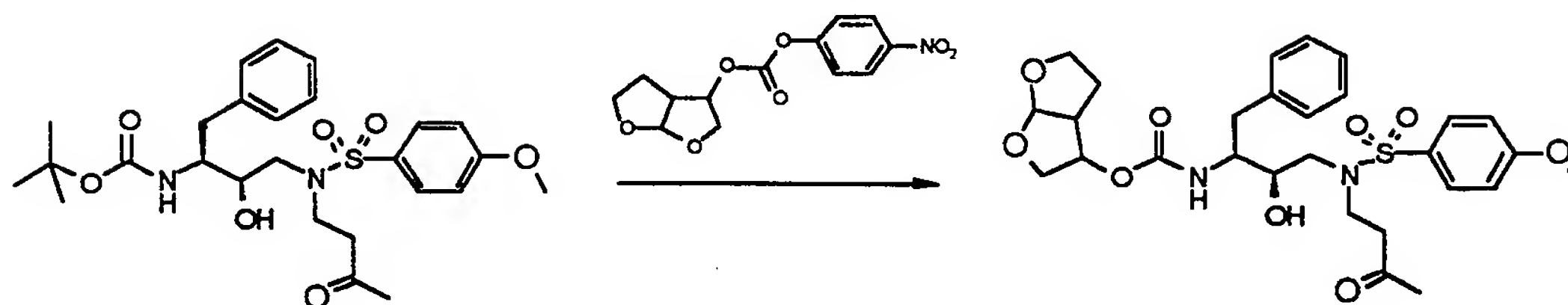
Step 2:



[(2R,3S) -3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino-(3-butanonyl (4-methoxyphenylsulfonyl)) butane].

[(2R,3S) -3-*tert*-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino-3-butanone) butane] (124 mg, 0.4 mmol) was combined with 4-methoxyphenylsulfonyl chloride (146 mg, 0.7 mmol) in DMF and cooled to 0 °C. This was followed by the addition of diisopropyl ethyl amine (186 uL, 1.1 mmol). The reaction was then warmed to room temperature and stirred for 15 hours. The reaction was quenched by the addition of 3 mLs of saturated NaHCO₃, followed by washing with sat. NaHCO₃, KHSO₃ and then brine. The organic phase was dried with MgSO₄ and the solvent was removed in vacuo. Purification by SiO₂ (1:3, EtOAc:Hex to 2:1 EtOAc:Hex). Isolated 0.038 g of the product as a colorless oil: 20.8 % yield, R_f=0.27 (1:1 EtOAc: Hex, /SiO₂), MS(ES) = 521 amu (M+1).

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Example (Compound 280)

{ (1S,2R) -3- [(4-methoxybenzenesulfonyl) - (3-butanonyl) - amino] -1-benzyl-2-hydroxy-propyl } -carbamic acid

5 (3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl ester.

Step 1:

[(2R,3S) -3-tert-Butoxycarbonylamino-2-hydroxy-4-phenyl-(1-amino-(3-butanonyl (4-methoxyphenylsulfonyl)) butane] (33.2 mg, 0.1 mmol) was dissolved in 1 mL of

10 CH₂Cl₂ followed by the addition of 1 mL of TFA. The reaction was stirred at room temperature for 15 minutes followed by concentration *in vacuo* to provide the desired deprotected amine. This material was used without further purification.

15 Step 2:

25.2 mg (0.03 mmol) of the material provided in the above deprotection step was then dissolved into DMF (1 mL) followed by the 3R,3aS,6aR)hexahydro-furo[2,3-b]furan-3-yl derived mixed carbonate (18.8 mg, 0.03

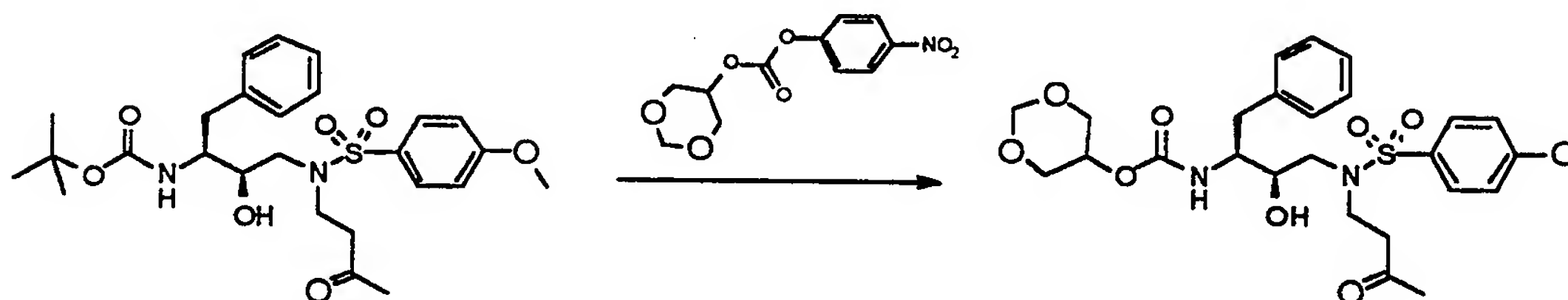
20 mmol). After cooling to 0 °C diisopropylethyl amine (33.3 uL, 0.2 mmol) was added and the reaction was stirred for 15 hours (warmed to room temperature overnight). The reaction mixture was then diluted with EtOAc washed with sat. NaHCO₃, KHSO₃ and then brine. The organic phase was

25 dried with MgSO₄ filtered and the solvent was removed *in vacuo*. Purification by Prep. TLC (SiO₂ 2:1, EtOAc:Hex).

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Isolated 0.0116 g of the product as a white foam: $R_f=0.16$ (2:1 EtOAc: Hex, /SiO₂), MS(ES) = 577 amu (M+1).

Example (Compound 281)

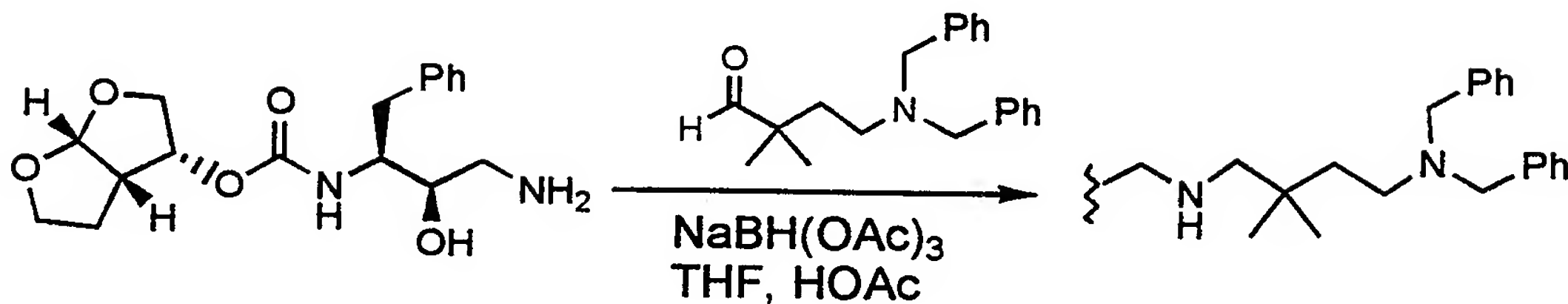


- 5 18.8 mg (0.03 mmol) of the material provided in
the above deprotection step (step 1 example 279) was then
dissolved into DMF (1 mL) followed by the 1,3-dioxan-5-yl
derived mixed carbonate (17.2 mg, 0.1 mmol). After
cooling to 0 °C diisopropylethyl amine (33.3 uL, 0.2 mmol)
10 was added and the reaction was stirred for 15 hours (
warmed to room temperature overnight). The reaction
mixture was then diluted with EtOAc washed with sat.
NaHCO₃, KHSO₃ and then brine. The organic phase was
dried with MgSO₄ and the solvent was removed *in vacuo*.
15 Purification by Prep. TLC (SiO₂ 2:1, EtOAc:Hex).
Isolated 0.0063 g of the product as a white foam: $R_f=0.29$
(2:1 EtOAc: Hex, /SiO₂), MS(ES) = 551 amu (M+1).

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Example (Compound 282)

Step 1:



5

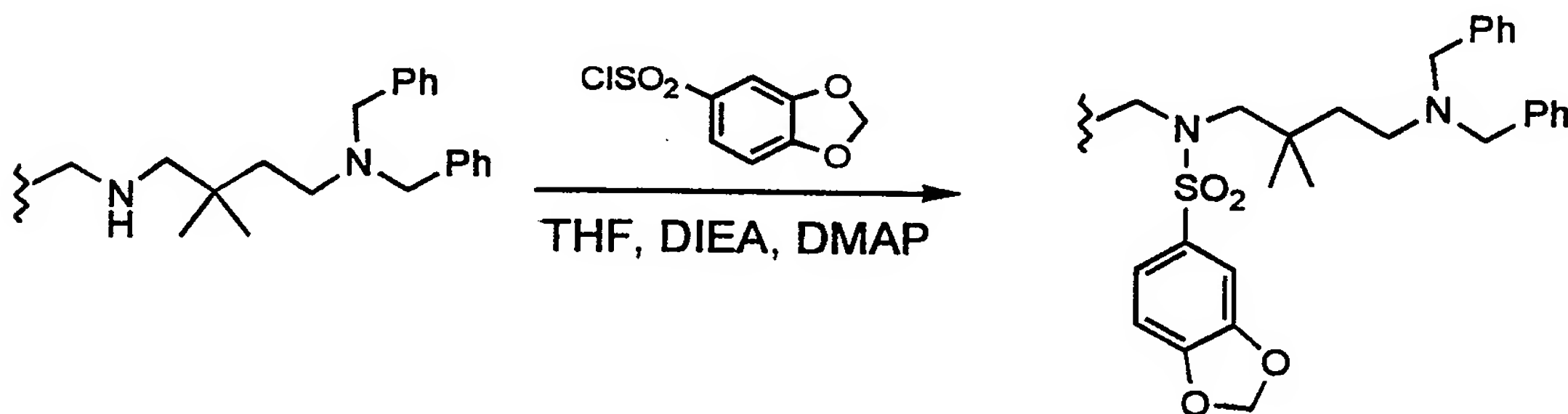
(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-[N,N-dibenzylamino]-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

According to example 257, step 5,

10 (3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-amino-1-benzyl-2-hydroxypropyl]carbamate was subjected to reductive alkylation with of 4-(N,N-dibenzylamino)-2,2-dimethylbutyraldehyde to afford the desired compound as thick oil in 70% yield. MS (ESI): 616 (M+H).

15

Step 2:



20 (3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl) (4-[N,N-dibenzylamino]-2,2-dimethylbutyl)amino]-1-benzyl-2-hydroxypropylcarbamate

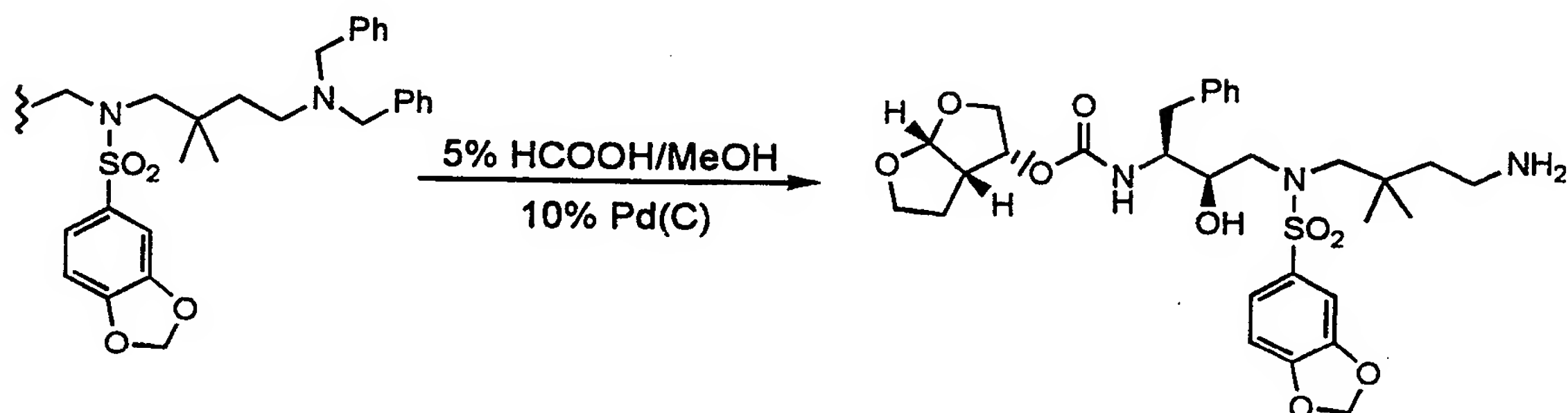
According to example 257, step 6,

(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-[N,N-dibenzylamino]-2,2-dimethylbutyl)amino]-

-271-

2-hydroxypropylcarbamate was subjected to sulfonylation to afford the desired sulfonamide as a white foam in 89% yield. MS (ESI): 800 (M+H).

5 Step 3:



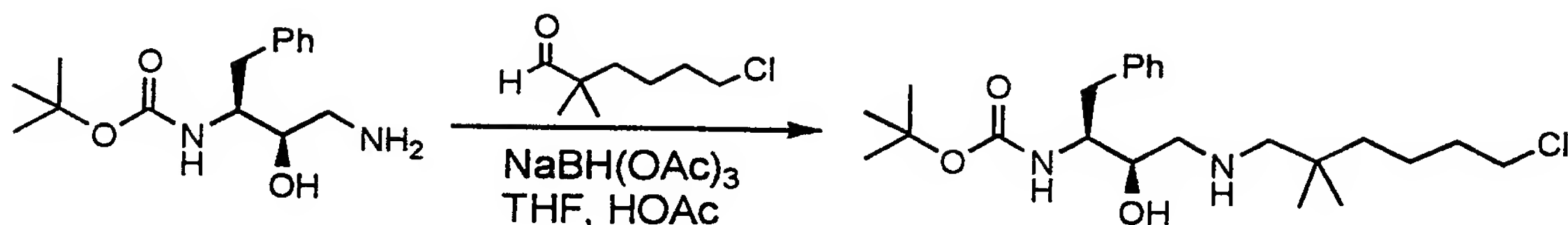
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 10 [(4-amino-2,2-dimethylbutyl) (1,3-benzodioxol-5-
 ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate
 According to example 217, step 3,
 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl) (4-[N,N-dibenzylamino]-
 15 2,2-dimethylbutyl) amino]-1-benzyl-2-ydroxypropylcarbamate
 was subjected to catalytic transfer hydrogenation to
 afford the desired amine as a white foam in 68% yield.
 NMR (CDCl₃): 7.34 (dd, 1H), 7.27-7.13 (m, 7H), 6.95 (d,
 1H), 6.15 (s, 2H), 5.81 (d, 1H), 5.61 (d, 1H), 5.01 (q,
 20 1H), 4.02 (t, 1H), 3.92-3.79 (3H), 3.67 (m, 2H), 3.20-
 2.96 (m, 4H), 2.91-2.60 (m, 5H), 2.45-1.70 (br, 2H),
 1.63-1.34 (m, 4H), 0.97 (s, 3H), 0.89 (s, 3H). MS (ESI):
 620 (M+H).

25

Example (Compound 283)

Step 1:

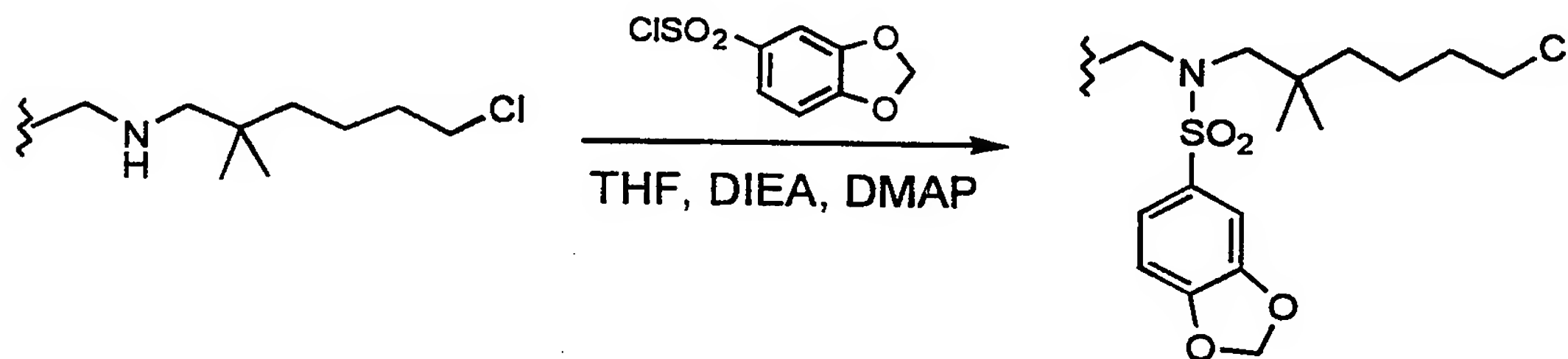
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5 **tert-butyl N-((1S,2R)-1-benzyl-3-[(6-chloro-2,2-dimethylhexyl)amino]-2-hydroxypropyl)carbamate**

According to example 257, step 5, tert-butyl N-[(1S,2R)-3-amino-1-benzyl-2-hydroxypropyl]carbamate was subjected to reductive alkylation with 6-chloro-2,2-dimethylhexanal to afford the desired compound as a viscous oil in
10 quantitative yield. MS (ESI): 428 (M+H).

Step 2:

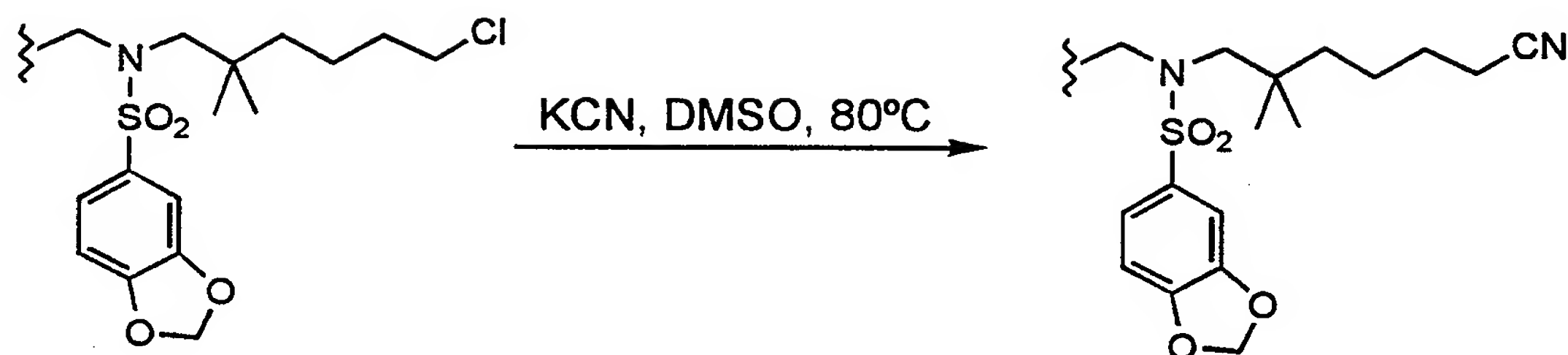


15 **tert-butyl N-((1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-chloro-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropyl)carbamate**

According to example 257, step 6, tert-butyl N-((1S,2R)-1-benzyl-3-[(6-chloro-2,2-dimethylhexyl)amino]-2-hydroxypropyl)carbamate was subjected to sulfonylation to afford the desired sulfonamide as a white foam in 70%
20 yield. MS (ESI): 633 (M+Na).

Step 3:

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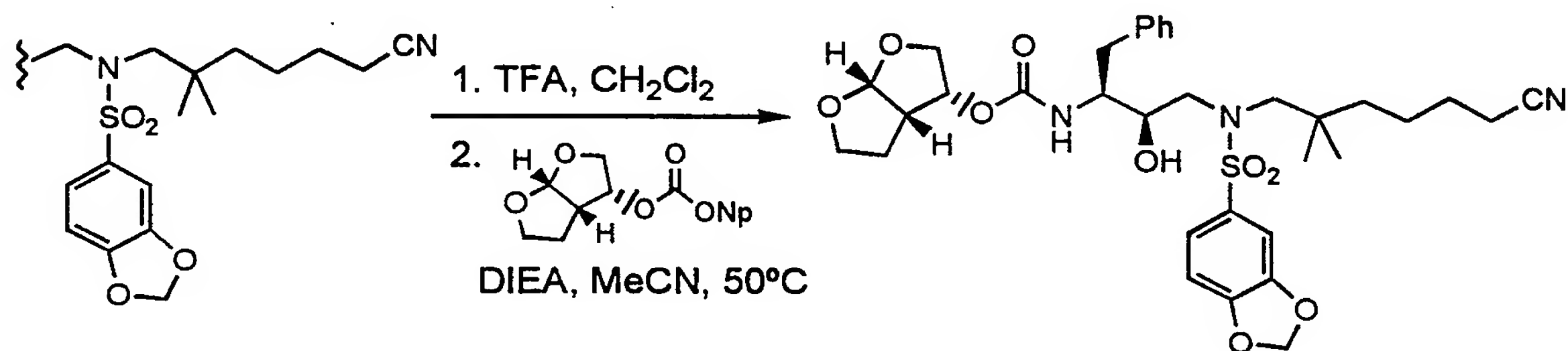


tert-butyl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-cyano-2,2-dimethylhexyl)amino]-1-benzyl-2-

5 hydroxypropylcarbamate

A solution of 200 mg (0.327 mmol) of tert-butyl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-chloro-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropylcarbamate and 43 mg (0.490 mmol) of KCN in 3 mL of DMSO was heated to 80°C with stirring. After 2 hours the solution was cooled to RT, diluted with water and the mixture extracted with CH₂Cl₂ (3x). The combined extracts were washed with brine (3x), dried over MgSO₄, and concentrated in vacuo to afford 196 mg (99%) of the desired compound as a light yellow foam. MS(ESI): 602 (M+H).

Step 4:



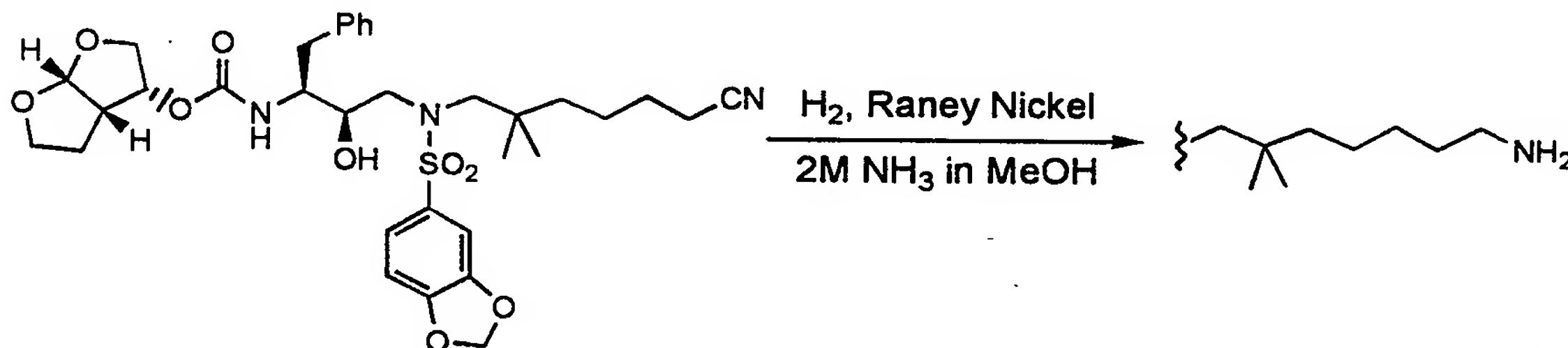
20 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-cyano-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropyl]carbamate

According to example 245, steps 4 and 5, tert-butyl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-cyano-2,2-

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dimethylhexyl) amino]-1-benzyl-2-hydroxypropylcarbamate was converted to the desired product in 66% overall yield. NMR (CDCl₃): 7.32 (dd, 1H), 7.30-7.12 (m, 6H), 6.89 (d, 1H), 6.07 (s, 2H), 5.61 (d, 1H), 4.97 (q, 1H), 4.88 (d, 1H), 4.02 (m, 1H), 3.92 (m, 2H), 3.81 (m, 2H), 3.65 (m, 2H), 3.17 (dd, 1H), 3.10-2.68 (m, 6H), 2.35 (t, 2H), 1.60 (m, 3H), 1.40 (m, 3H), 1.29 (m, 2H), 0.90 (d, 6H). MS (ESI): 658 (M+H).

10 Step 5:



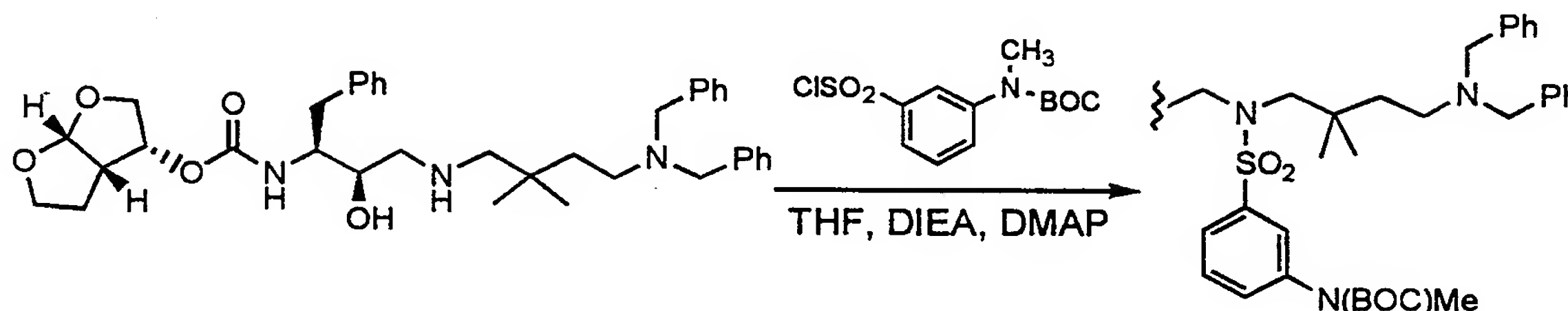
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
15 [(1,3-benzodioxol-5-ylsulfonyl)(7-amino-2,2-
dimethylheptyl) amino]-1-benzyl-2-hydroxypropyl]carbamate
According to example 245, step 6,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(6-cyano-2,2-
20 dimethylhexyl) amino]-1-benzyl-2-hydroxypropyl]carbamate
was subjected to Raney nickel hydrogenation to afford the
desired primary amine as a white foam in 89% yield. H1-
NMR (CDCl₃): 7.40 (d, 1H), 7.28-7.16 (m, 6H), 6.86 (d,
1H), 6.12 (s, 2H), 5.75 (d, 1H), 5.62 (m, 1H), 5.03 (m,
25 1H), 4.60-3.70 (br, 2H), 4.21-3.61 (m, 6H), 3.25-2.71 (m,
10H), 1.85-1.24 (m, 10H), 0.95 (d, 6H).

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Example (Compound 284)

Step 1:



5

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-[(tert-butoxycarbonyl)(methyl)amino]phenylsulfonyl)(4-[N,N-dibenzylamino]-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

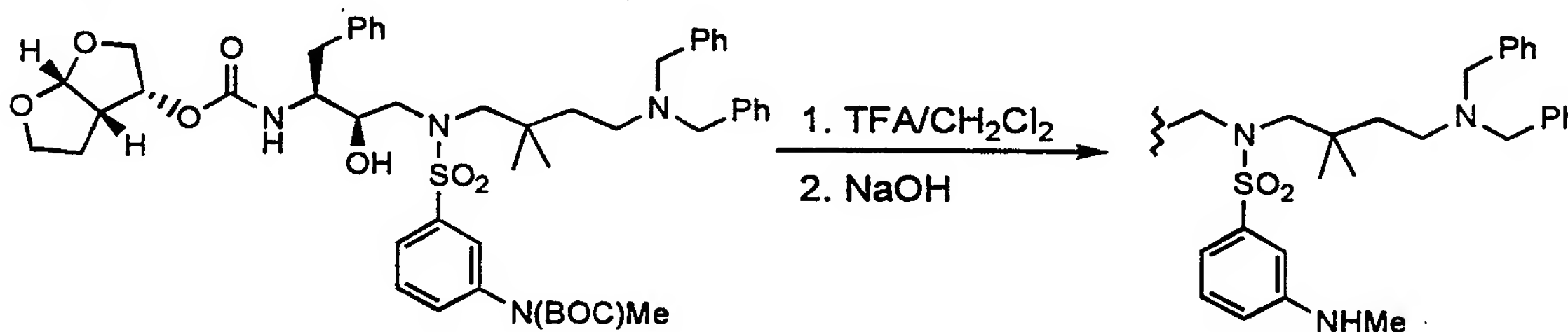
10

According to example 258, step 1,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-[N,N-dibenzylamino]-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate was

15 subjected to sulfonylation to give the desired product as a light yellow foam in 54% yield. MS(ESI): 886(M+H).

Step 2:



20

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-(methylamino)phenylsulfonyl)(4-[N,N-dibenzylamino]-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

25

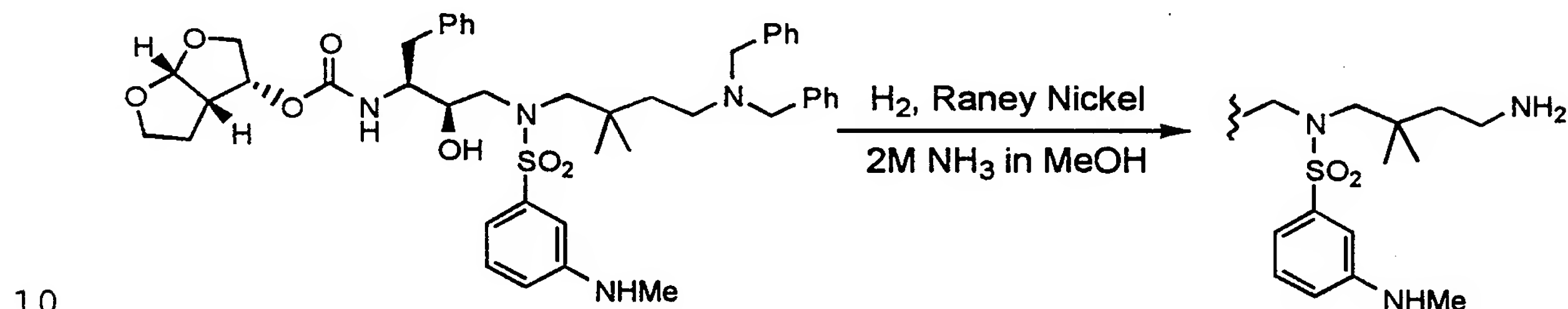
-276-

According to example 258, step 2,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-[(tert-butoxycarbonyl)(methyl)amino]phenylsulfonyl)(4-[N,N-dibenzylamino]-2,2-

5 dimethylbutyl)amino]-2-hydroxypropylcarbamate was subjected to TFA deprotection to afford the desired product in 96% yield. MS(ESI): 786(M+H).

Step 3:



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-(methylamino)phenylsulfonyl)(4-amino-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

15 According to example 245, step 6,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-(methylamino)phenylsulfonyl)(4-[N,N-dibenzylamino]-2,2-dimethylbutyl)amino]-2-

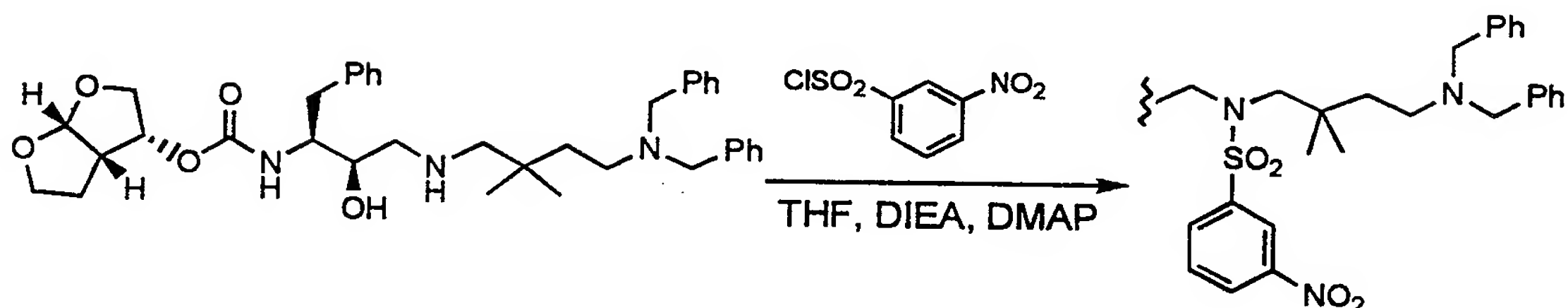
20 hydroxypropylcarbamate was subjected to Raney nickel reduction to afford the desired primary amine as a white foam in 45% yield. NMR (CDCl₃): 7.36-7.17 (m, 6H), 7.02 (d, 1H), 6.95 (s, 1H), 6.76 (dd, 1H), 5.78 (d, 1H), 5.60 (d, 1H), 4.99 (q, 1H), 4.11 (m, 1H), 4.02 (t, 1H), 3.91-3.78 (m, 3H), 3.73-3.59 (m, 2H), 3.25-2.99 (m, 5H), 2.83 (m, 5H), 2.78-2.59 (m, 3H), 2.60-1.90 (br, 2H), 1.63-1.33 (m, 4H), 0.98 (s, 3H), 0.89 (s, 3H). MS(ESI): 605(M+H).

25

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Example (Compound 285)

Step 1:



5

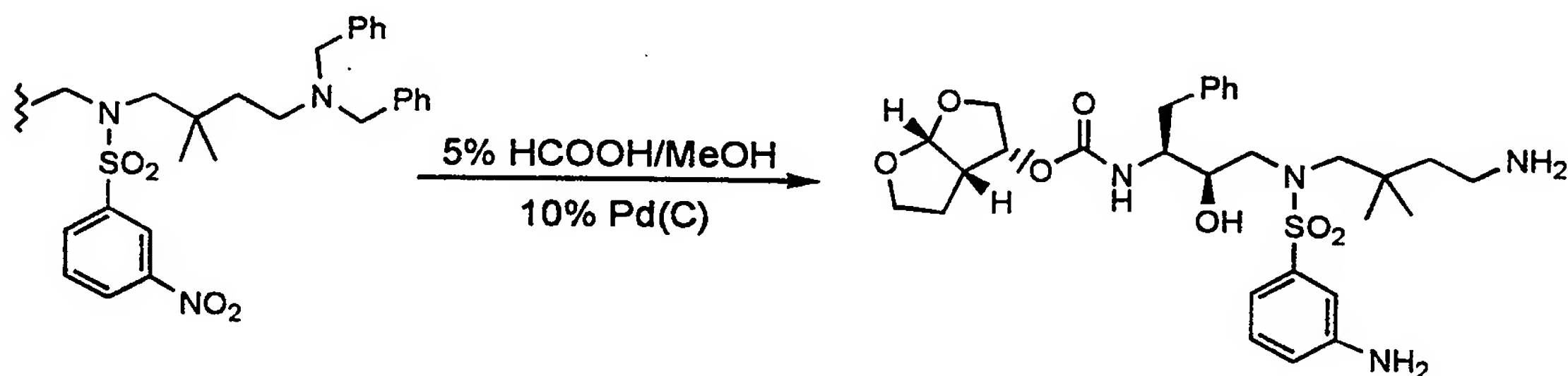
(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-nitrophenylsulfonyl) (4-[N,N-dibenzylamino]-2,2-dimethylbutyl) amino]-2-hydroxypropylcarbamate

According to example 257, step 6,

10 (3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-[N,N-dibenzylamino]-2,2-dimethylbutyl) amino]-2-hydroxypropylcarbamate was subjected to sulfonylation with 3-nitrobenzenesulfonyl chloride to afford the desired sulfonamide as a white foam in 90% yield.

15 MS (ESI): 801 (M+H).

Step 2:



20 (3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-aminophenylsulfonyl) (4-amino-2,2-dimethylbutyl) amino]-2-hydroxypropylcarbamate

According to example 217, step 3,

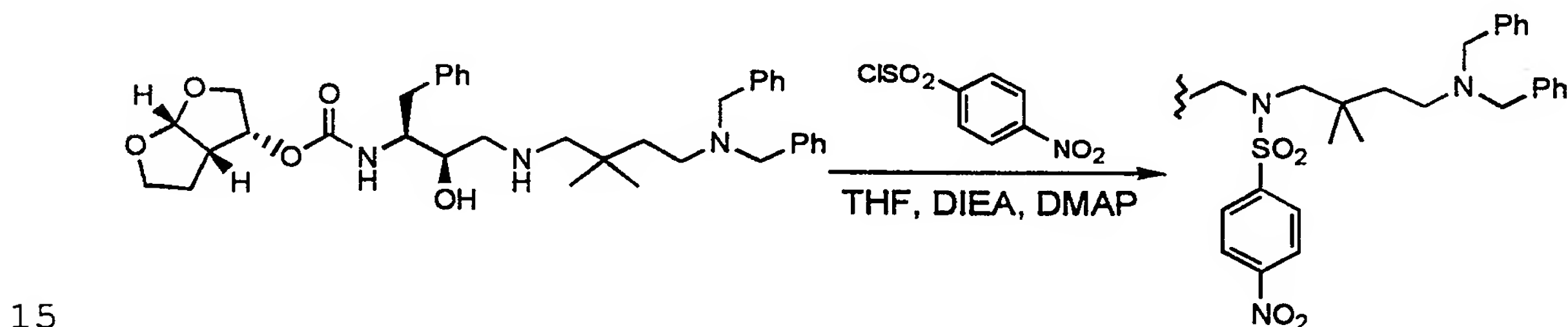
(3R,3aS,6aR) hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-

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benzyl-3-[(3-nitrophenylsulfonyl) (4-[N,N-dibenzylamino]-
2,2-dimethylbutyl) amino]-2-hydroxypropylcarbamate was
subjected to catalytic transfer hydrogenation to afford
the desired primary amine as a white foam in 63% yield.
5 NMR (CDCl₃): 7.37-7.19 (m, 6H), 7.14 (m, 2H), 6.88 (dd,
1H), 5.84 (d, 1H), 5.65 (d, 1H), 5.04 (q, 1H), 4.10 (m,
3H), 3.99-3.81 (m, 3H), 3.80-3.64 (m, 2H), 3.32-3.01 (m,
5H), 2.91 (m, 2H), 2.82-2.65 (m, 3H), 2.60-2.00 (br, 2H),
1.70-1.39 (m, 4H), 1.04 (s, 3H), 0.95 (s, 3H). MS (ESI):
10 591 (M+H).

Example (Compound 286)

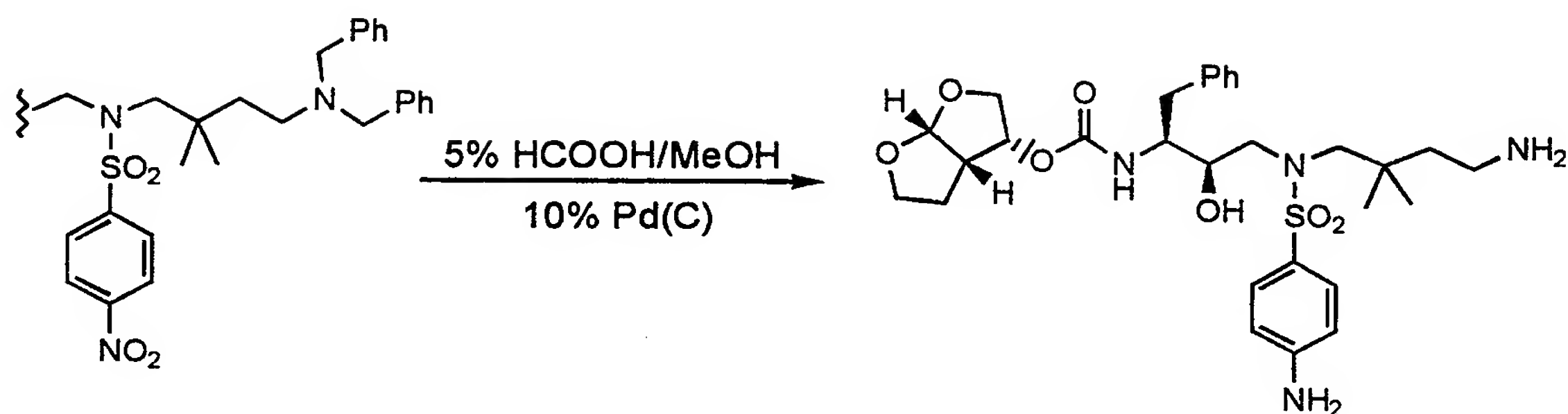
Step 1:



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-
benzyl-3-[(4-nitrophenylsulfonyl) (4-[N,N-dibenzylamino]-
2,2-dimethylbutyl) amino]-2-hydroxypropylcarbamate
20 According to example 257, step 6,
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-
benzyl-3-[(4-[N,N-dibenzylamino]-2,2-dimethylbutyl) amino]-
2-hydroxypropylcarbamate was subjected to sulfonylation
25 with 4-nitrobenzenesulfonyl chloride to afford the
desired sulfonamide as a light yellow foam in 86% yield.
MS (ESI): 801 (M+H).

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Step 2:



5 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-aminophenylsulfonyl)(4-amino-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

According to example 217, step 3,

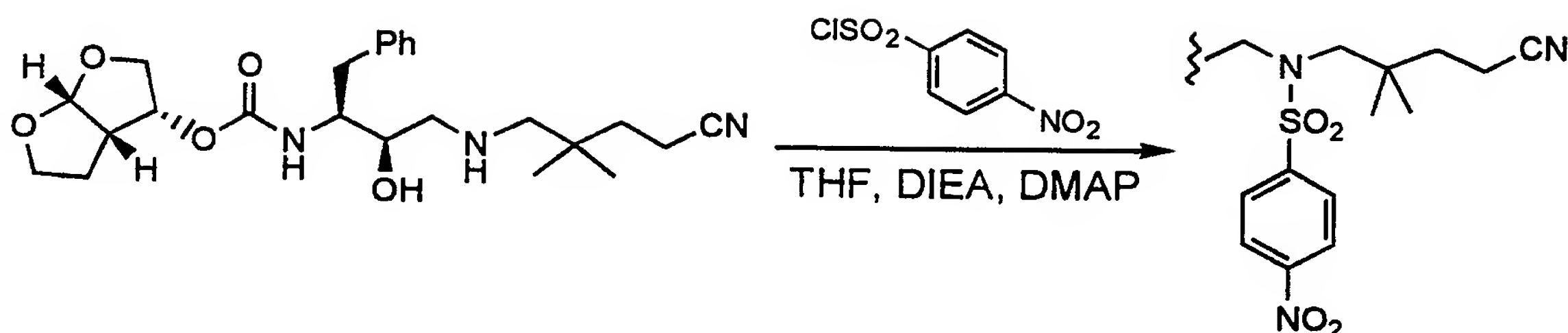
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-nitrophenylsulfonyl)(4-[N,N-dibenzylamino]-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate was subjected to catalytic transfer hydrogenation to afford the desired primary amine as a white foam in 63% yield. NMR (CDCl₃): 7.55 (d, 2H), 7.28-7.15 (m, 5H), 6.67 (d, 2H), 5.77 (d, 1H), 5.60 (d, 1H), 4.99 (q, 1H), 4.20 (s, 2H), 4.02 (t, 1H), 3.92-3.78 (m, 3H), 3.71-3.61 (m, 2H), 3.20-3.02 (m, 4H), 2.96 (d, 1H), 2.85 (q, 1H), 2.80-2.60 (m, 4H), 2.60-1.80 (br, 2H), 1.63-1.35 (m, 4H), 0.96 (s, 3H), 0.89 (s, 3H). MS(ESI): 591(M+H).

20

Example (Compound 287)

Step 1:

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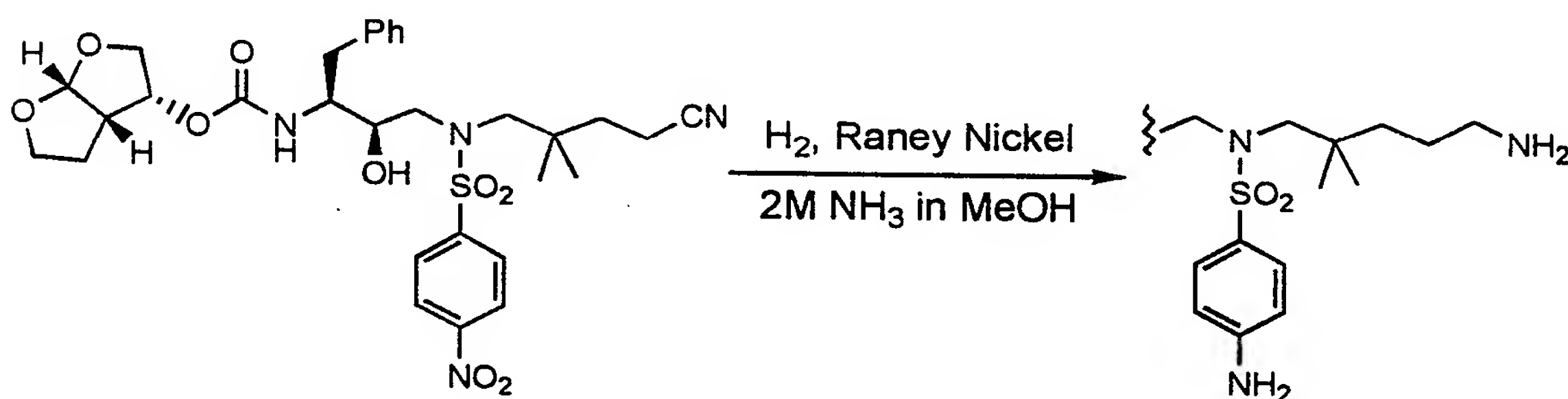


(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-nitrophenylsulfonyl)(4-cyano-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

According to example 257, step 6,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-cyano-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate (example 257, step 5) was subjected to sulfonylation with 4-nitrobenzenesulfonyl chloride to afford the desired sulfonamide as a light yellow foam in 82% yield. MS(ESI): 653(M+Na).

Step 2:



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-aminophenylsulfonyl)(5-amino-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate

According to example 245, step 6,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-nitrophenylsulfonyl)(4-cyano-2,2-

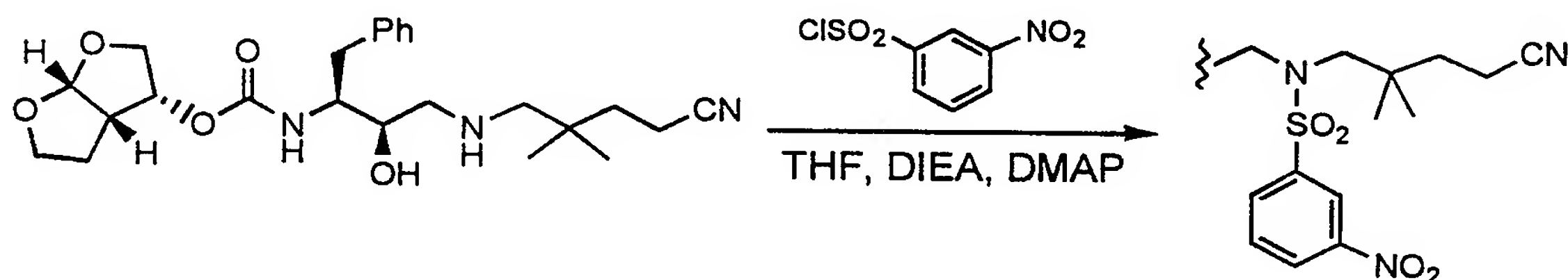
-281-

dimethylbutyl)amino]-2-hydroxypropylcarbamate was subjected to Raney nickel hydrogenation to afford the desired amine as a white foam in 79% yield. NMR (CDCl₃): 7.60 (d, 2H), 7.38-7.18 (m, 5H), 7.00 (d, 1H), 6.72 (d, 2H), 5.65 (d, 1H), 5.05 (q, 1H), 4.21 (s, 2H), 4.07-3.78 (m, 4H), 3.76-3.62 (m, 2H), 3.36-3.07 (m, 3H), 3.05-2.72 (m, 5H), 2.61 (m, 2H), 2.55-1.60 (br, 2H), 1.80-1.28 (m, 6H), 1.02 (s, 3H), 0.90 (s, 3H). MS(ESI): 605 (M+H).

Example (Compound 288)

10

Step 1:



15 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-nitrophenylsulfonyl)(4-cyano-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

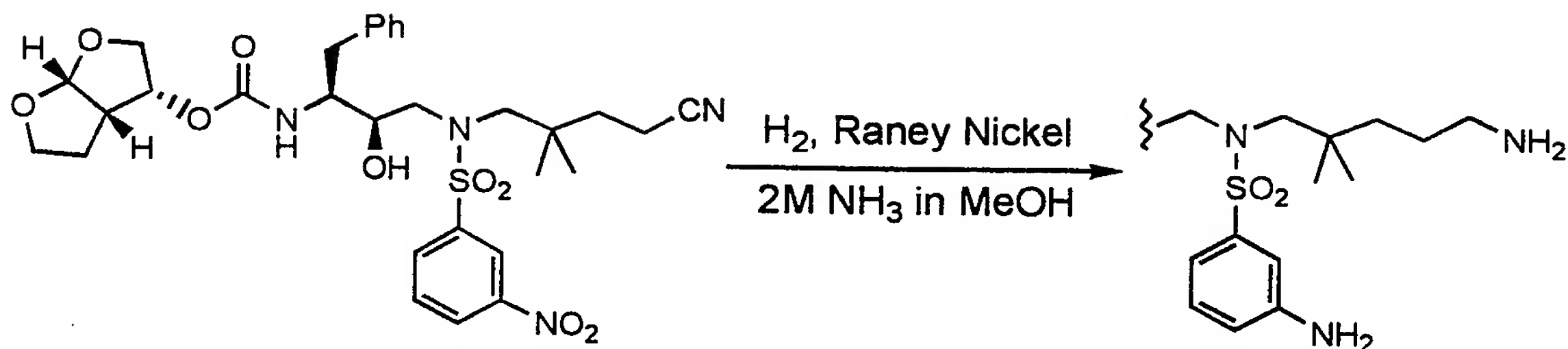
According to example 257, step 6,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-cyano-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate (example 257, step 5) was subjected to sulfonylation with 3-nitrobenzenesulfonyl chloride to afford the desired sulfonamide as a white foam in 80% yield. MS(ESI): 653 (M+Na).

25

Step 2:

-282-



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-aminophenylsulfonyl)(5-amino-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate

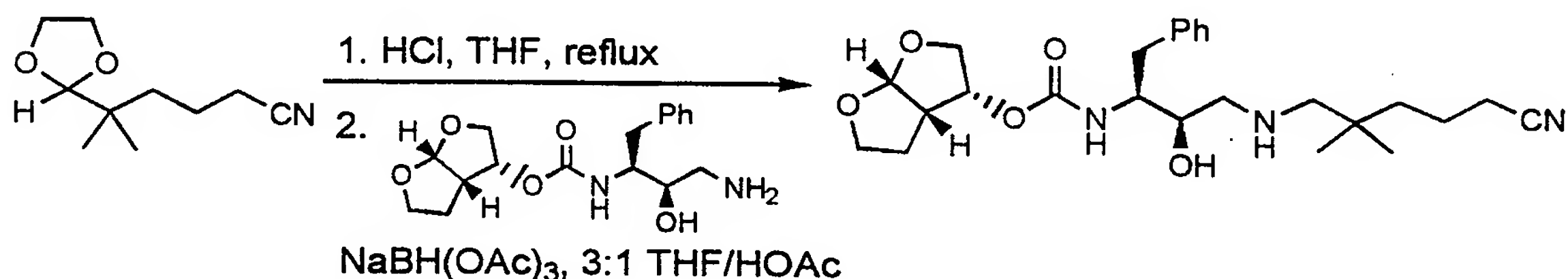
According to example 245, step 6,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-nitrophenylsulfonyl)(4-cyano-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate was subjected to Raney nickel hydrogenation to afford the desired amine as a white foam in 91% yield. NMR (CDCl₃): 7.37-7.08 (m, 8H), 7.00 (d, 1H), 6.89 (dd, 1H), 5.64 (d, 1H), 5.05 (q, 1H), 4.13-3.79 (m, 6H), 3.70 (m, 2H), 3.36-3.00 (m, 4H), 2.98-2.57 (m, 6H), 2.50-1.70 (br, 2H), 1.75-1.23 (m, 6H), 1.02 (s, 3H), 0.91 (s, 3H). MS(ESI): 605 (M+H).

Example (Compound 289)

20

Step 1:

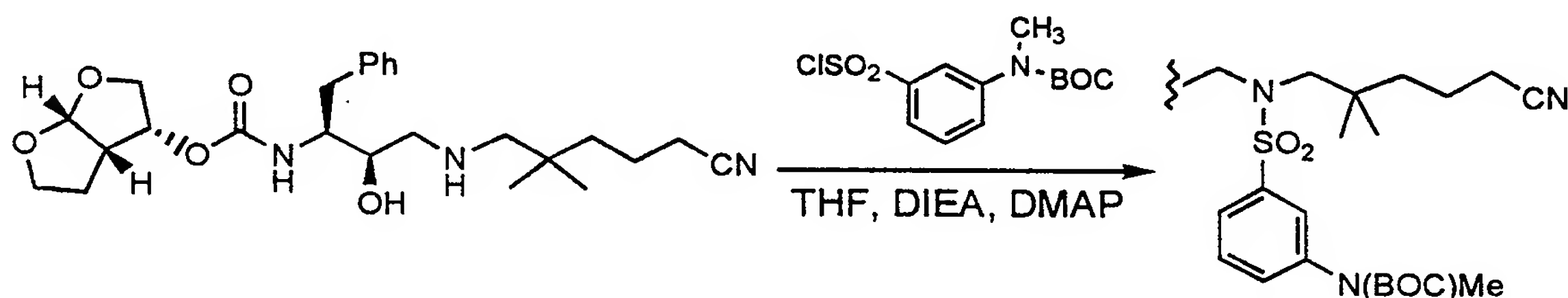


-283-

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate

According to example 18, steps 1 and 2, 4-(1,3-dioxolan-2-yl)-4-methylpentanenitrile was subjected to acidic hydrolysis followed by reductive amination with (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-amino-1-benzyl-2-hydroxypropyl]carbamate (example 257, step 4) except that 3:1 THF/glacial acetic acid was used as the solvent in the second step. The crude product was purified by flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to give the desired compound as a thick semi-solid in 58% yield. MS(ESI): 460 (M+H).

Step 2:

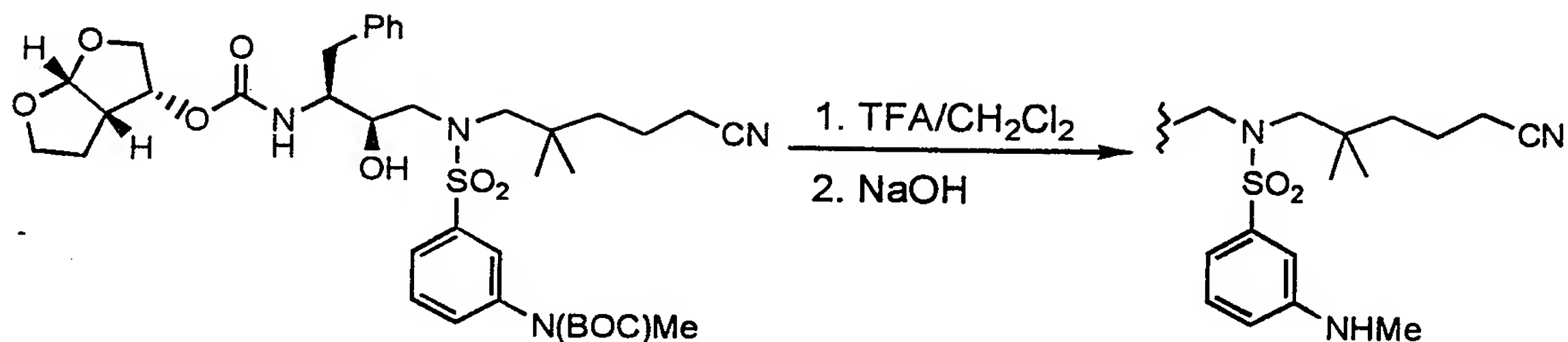


(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-[(tert-butoxycarbonyl)(methyl)amino]phenylsulfonyl)(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate

According to example 258, step 1, (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate was subjected to sulfonylation to give the desired product as a white foam in 35% yield. MS(ESI): 751 (M+Na).

-284-

Step 3:



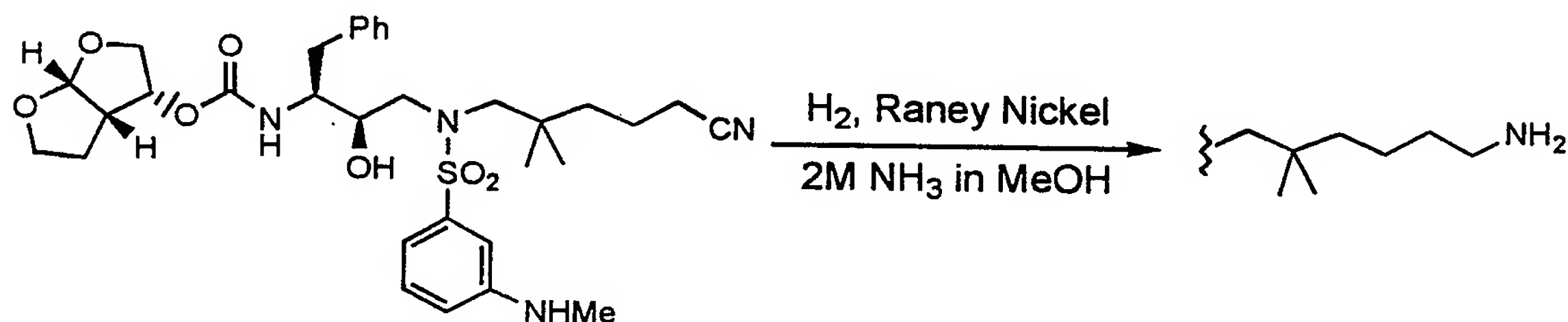
- 5 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-(methylanino)phenylsulfonyl)(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate

According to example 258, step 2,

- 10 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-[(tert-butoxycarbonyl)(methyl)amino]phenylsulfonyl)(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate was subjected to TFA deprotection to afford the desired product in 94% yield. MS(ESI): 629 (M+H).

15

Step 4:



- 20 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-(methylanino)phenylsulfonyl)(6-amino-2,2-dimethylhexyl)amino]-2-hydroxypropylcarbamate

According to example 245, step 6,

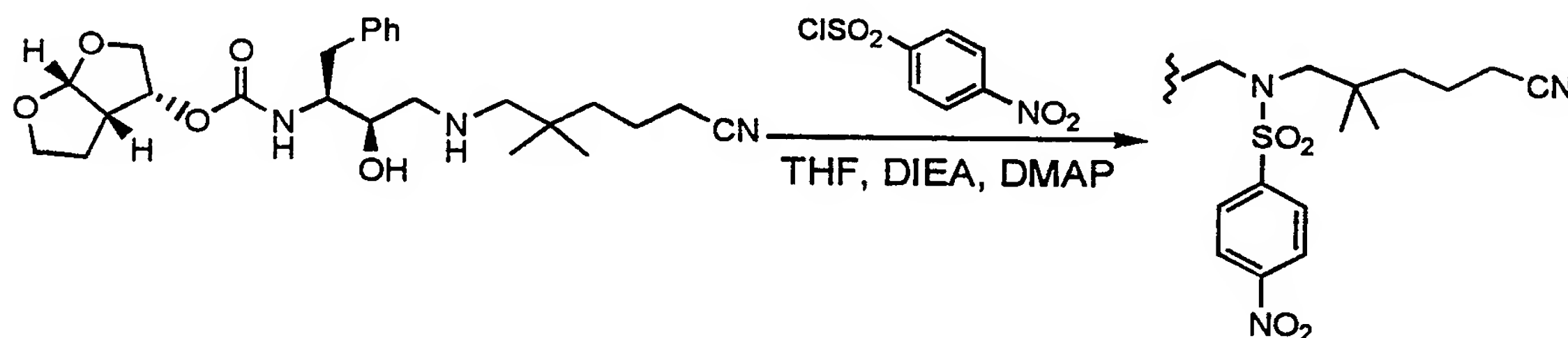
- 25 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-(methylanino)phenylsulfonyl)(5-cyano-2,2-

-285-

dimethylpentyl)amino]-2-hydroxypropylcarbamate was subjected to Raney nickel hydrogenation to afford the desired primary amine as a white foam in 79% yield. ¹H-NMR (CDCl₃): 7.37-7.17 (m, 6H), 7.09 (d, 1H), 7.01 (s, 1H), 6.80 (dd, 1H), 5.65 (d, 1H), 5.45 (d, 1H), 5.02 (q, 1H), 4.25 (br s, 1H), 4.06 (t, 1H), 4.01-3.63 (m, 5H), 3.33-2.40 (m, 15H), 1.72-1.27 (m, 8H), 0.96 (d, 6H). MS (ESI): 633 (M+H).

Example (Compound 290)

Step 1:



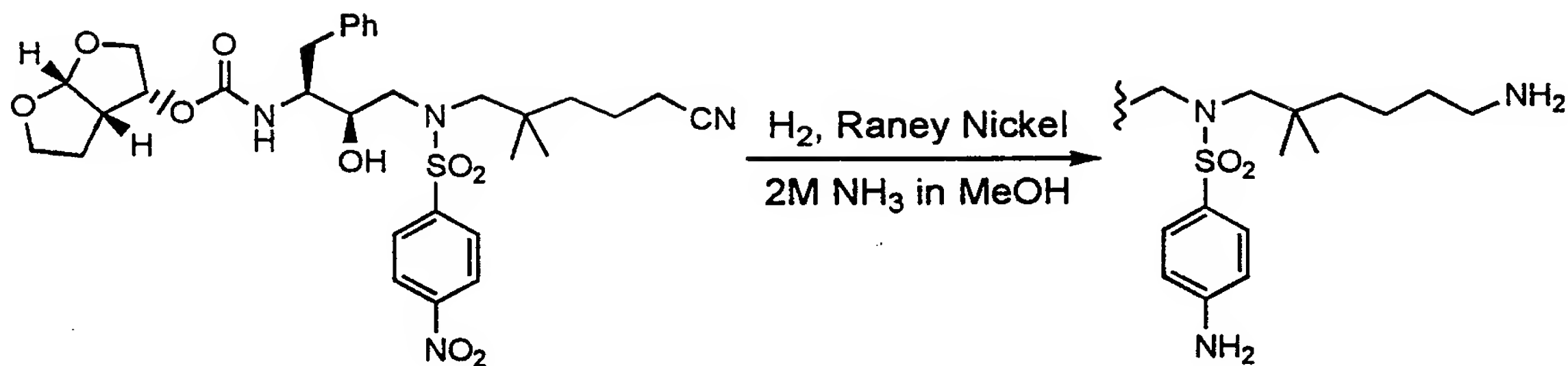
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-nitrophenylsulfonyl)(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate

According to example 257, step 6,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate (example 289, step 1) was subjected to sulfonylation with 4-nitrobenzenesulfonyl chloride to afford the desired sulfonamide as a light yellow foam in 87% yield. MS (ESI): 667 (M+Na).

Step 2:

-286-



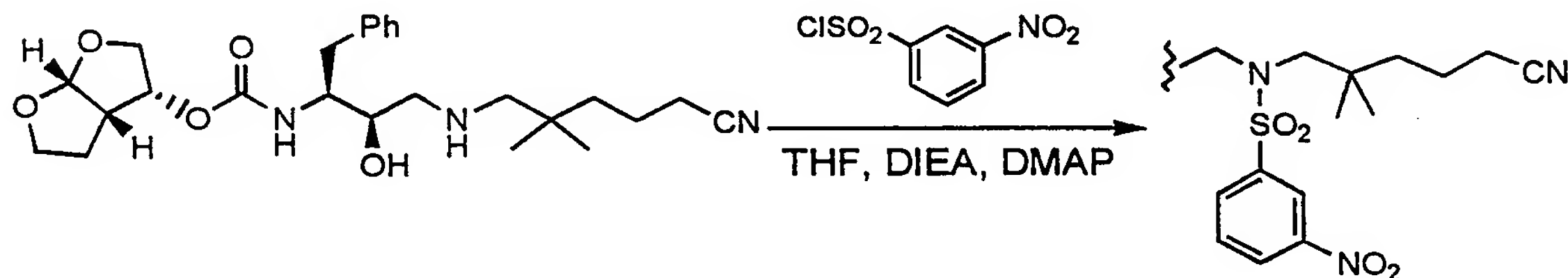
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-aminophenylsulfonyl)(6-amino-2,2-dimethylhexyl)amino]-2-hydroxypropylcarbamate

According to example 245, step 6,

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(4-nitrophenylsulfonyl)(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate was subjected to Raney nickel hydrogenation to afford the desired primary amine as a white foam in 79% yield. ¹H-NMR (CDCl₃): 7.60 (d, 2H), 7.33-7.18 (m, 5H), 6.75 (d, 2H), 5.66 (s, 1H), 5.49 (d, 1H), 5.02 (q, 1H), 4.32 (s, 2H), 4.09 (t, 1H), 4.01-3.65 (m, 5H), 3.27-2.50 (m, 12H), 1.70-1.23 (m, 8H), 0.95 (d, 6H). MS (ESI): 619 (M+H).

Example (Compound 291)

Step 1:



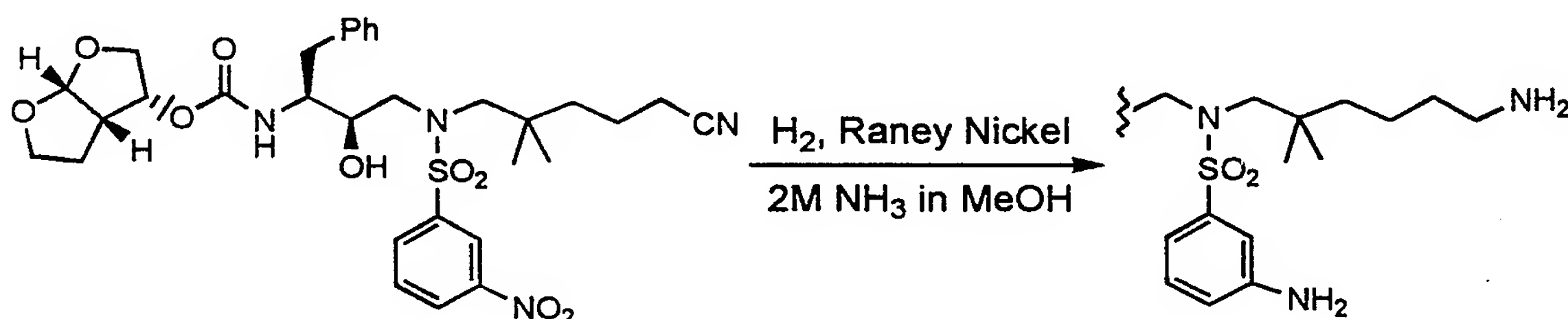
-287-

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-nitrophenylsulfonyl)(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate

According to example 257, step 6,

5 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate (example 289, step 1) was subjected to sulfonylation with 3-nitrobenzenesulfonyl chloride to afford the desired sulfonamide as a white
10 foam in 90% yield. MS(ESI): 667(M+Na).

Step 2:



15

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-aminophenylsulfonyl)(6-amino-2,2-dimethylhexyl)amino]-2-hydroxypropylcarbamate

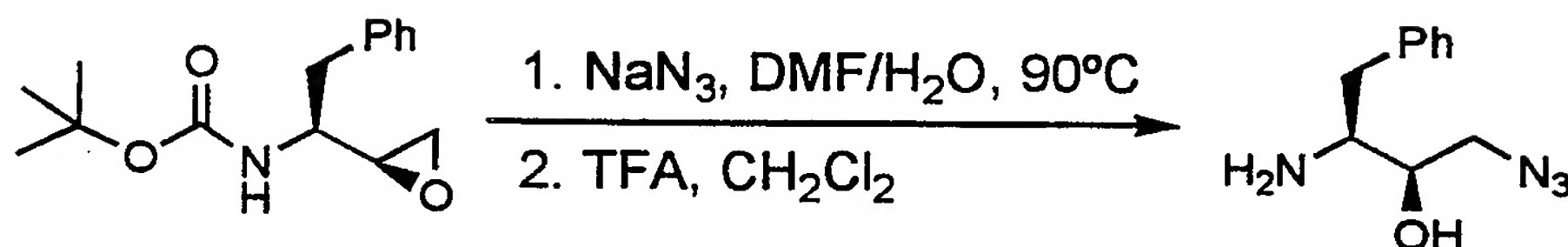
According to example 245, step 6,

20 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(3-nitrophenylsulfonyl)(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate was subjected to Raney nickel hydrogenation to afford the desired primary amine as a white foam in 78% yield. H1-
25 NMR (CDCl₃): 7.36-7.19 (m, 6H), 7.14 (m, 2H), 6.88 (dd, 1H), 5.66 (d, 1H), 5.46 (d, 1H), 5.04 (q, 1H), 4.30-4.01 (m, 3H), 4.00-3.65 (m, 5H), 3.33-2.40 (m, 12H), 1.72-1.25 (m, 8H), 0.95 (d, 6H). MS(ESI): 619(M+H).

-288-

Example (Compound 292)

Step 1:



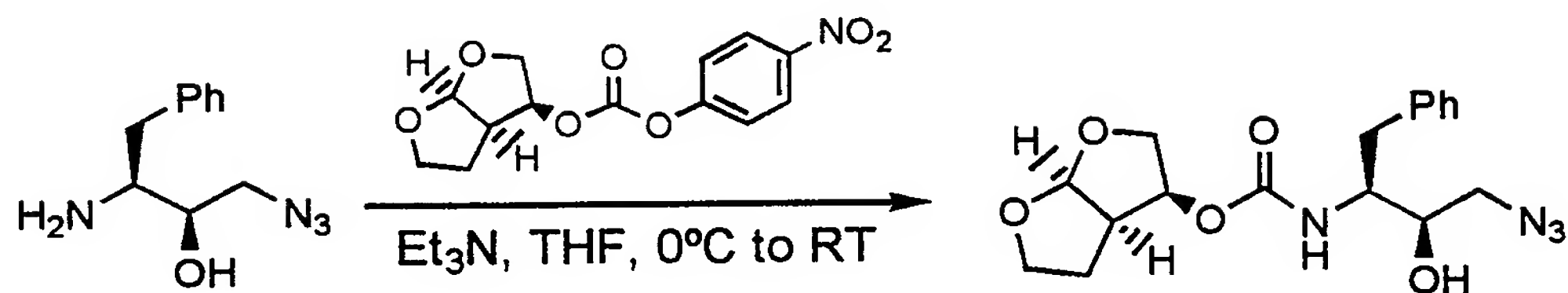
5

[(3S,2R)-3-amino-2-hydroxy-4-phenylbutyl]azide

A solution of 53 g (200 mMol) of tert-butyl N-((1S)-1-
[(2S)oxiran-2-yl]-2-phenylethylcarbamate and 20 g of
sodium azide (300 mMol) in 500 mL of DMF and 25 mL of
10 water was heated to 90°C for 3 hours. The mixture was
cooled and the product was precipitated by the addition
of 1 L of water. The solid was collected, re-dissolved
in ethyl acetate and extracted with 1 N hydrochloric acid
followed by saturated sodium bicarbonate. Drying over
15 magnesium sulfate and evaporation of the volatiles
afforded 31 g of the desired azide (m/e: 307 (M+H)) as a
white solid which was dissolved in 200 mL of
dichloromethane and treated with 200 mL of
trifluoroacetic acid. The resulting solution was stirred
20 at room temperature for 3 hours and the volatiles were
removed in vacuo. The residue was partitioned between
ethyl acetate and saturated sodium bicarbonate solution.
The organic layer was dried and evaporated to give the
free amine as a colorless syrup which solidified on
25 standing. MS(ESI): 207 (M+H).

Step 2:

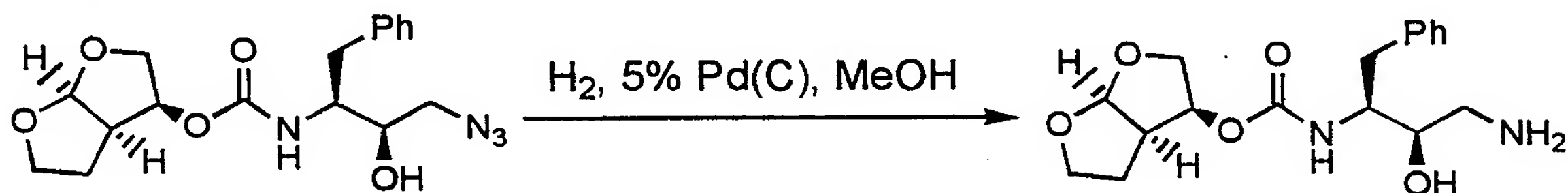
-289-



(3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-azido-1-benzyl-2-hydroxypropyl]carbamate

- 5 A solution of 1.91 g (9.26 mmol) of [(3S,2R)-3-amino-2-hydroxy-4-phenylbutyl]azide and 2.28 g (7.72 mmol) of (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl (4-nitrophenyl)carbonate in 40 mL of THF was treated with 2.20 mL (15.5 mmol) of triethylamine. The resulting
- 10 solution was stirred at RT. After 3 days the solution was concentrated in vacuo and the residue dissolved in CH₂Cl₂. The solution was washed with 5% aqueous citric acid (3x), 1M aqueous NaOH (3x) and then water (2x). The solution was dried over MgSO₄ and concentrated to afford
- 15 2.20 g (78%) of the desired carbamate as a light tan solid. MS(ESI): 385(M+H). The nitrophenylcarbonate used in the above procedure was prepared by treating (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-ol (Ghosh, Arun K.; Chen, Yan. Tetrahedron Lett. (1995), 36(4), 505-8)
- 20 with 1.05 equiv. of 4-nitrophenyl chloroformate in CH₂Cl₂ in the presence of 1.05 equiv. of pyridine.

Step 3:



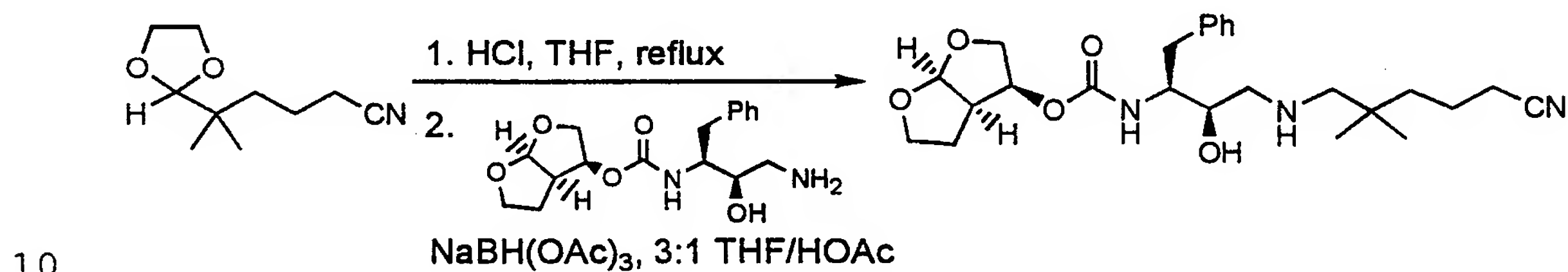
25

(3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-amino-1-benzyl-2-hydroxypropyl]carbamate

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According to example 257, step 4,
 (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-azido-1-benzyl-2-hydroxypropyl]carbamate was subjected to catalytic hydrogenation. The crude product was purified
 5 by flash chromatography (SiO₂, CH₂Cl₂/2M NH₃ in MeOH) to afford the desired amine as a light yellow solid in 77% yield. MS(ESI): 337(M+H).

Step 4:

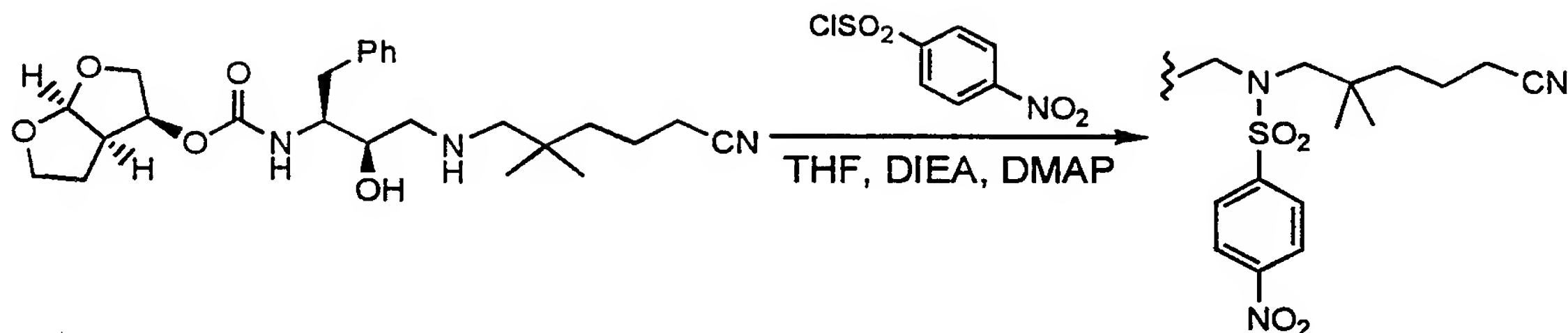


(3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl)amino]-2-hydroxypropylcarbamate
 15

According to example 18, steps 1 and 2, 4-(1,3-dioxolan-2-yl)-4-methylpentanenitrile was subjected to acidic hydrolysis followed by reductive amination with
 (3S,3aR,6aS)hexahydrofuro[2,3-b]furan-3-yl N-[(1S,2R)-3-amino-1-benzyl-2-hydroxypropyl]carbamate except that 3:1
 20 THF/glacial acetic acid was used as the solvent in the second step. The crude product was purified by flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to give the desired compound as a thick transparent oil in 52%
 25 yield. MS(ESI): 460(M+H).

Step 5:

-291-



(3S, 3aR, 6aS) hexahydrofuro[2,3-b]furan-3-yl N-(1S, 2R)-1-benzyl-3-[(4-nitrophenylsulfonyl) (5-cyano-2,2-dimethylpentyl) amino]-2-hydroxypropylcarbamate

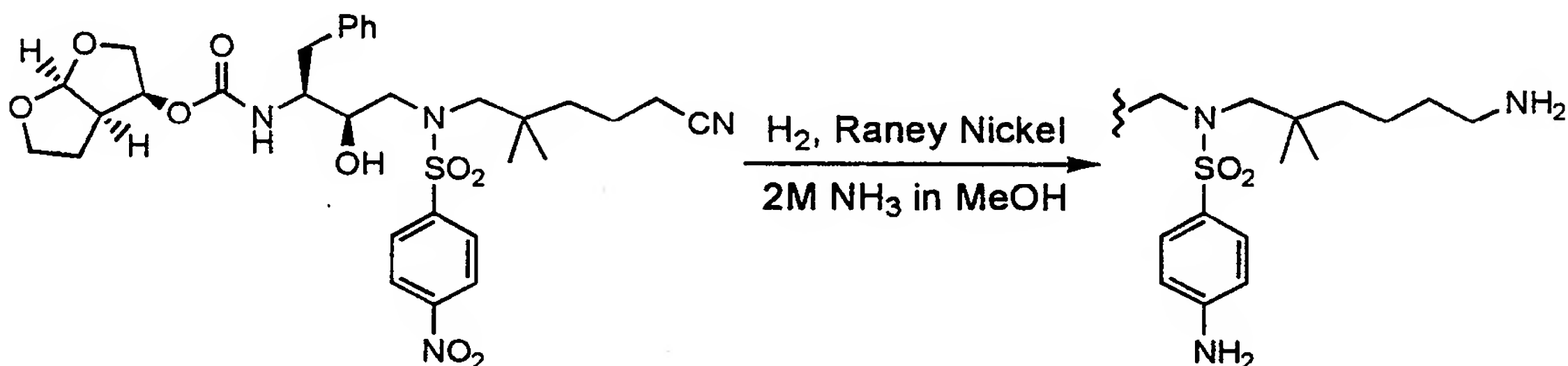
According to example 257, step 6,

(3S, 3aR, 6aS) hexahydrofuro[2,3-b]furan-3-yl N-(1S, 2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl) amino]-2-

hydroxypropylcarbamate was subjected to sulfonylation with 4-nitrobenzenesulfonyl chloride to afford the desired sulfonamide as a white foam in 93% yield.

MS (ESI): 667 (M+Na).

Step 6:



(3S, 3aR, 6aS) hexahydrofuro[2,3-b]furan-3-yl N-(1S, 2R)-1-benzyl-3-[(4-aminophenylsulfonyl) (6-amino-2,2-dimethylhexyl) amino]-2-hydroxypropylcarbamate

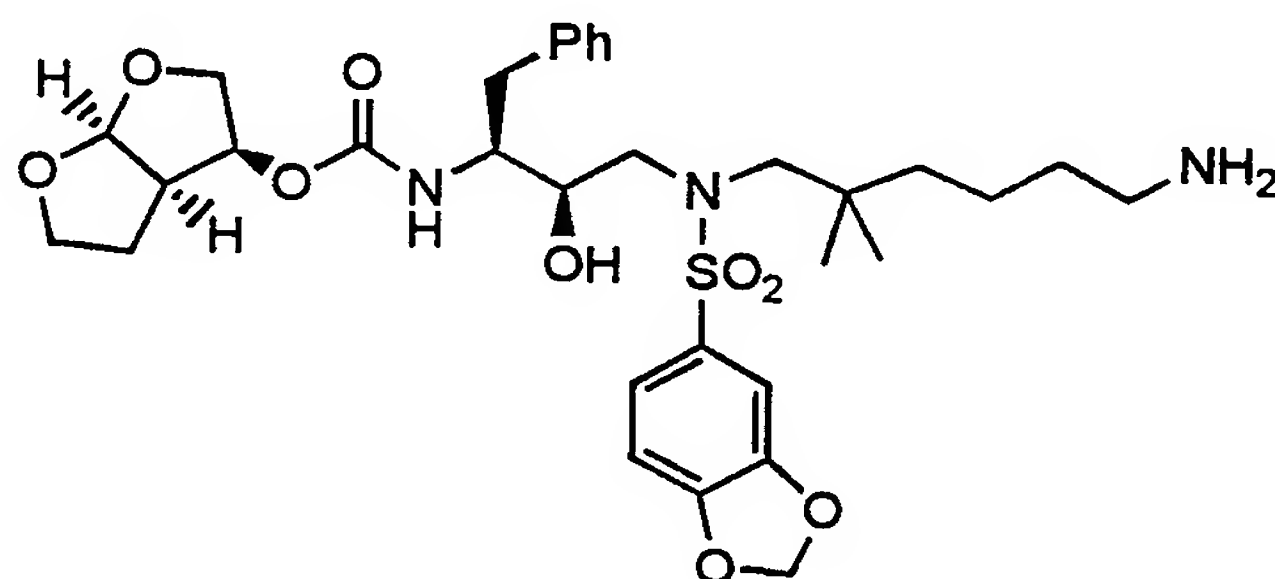
According to example 245, step 6,

(3S, 3aR, 6aS) hexahydrofuro[2,3-b]furan-3-yl N-(1S, 2R)-1-benzyl-3-[(4-nitrophenylsulfonyl) (5-cyano-2,2-

-292-

dimethylpentyl) amino]-2-hydroxypropylcarbamate was subjected to Raney nickel hydrogenation to afford the desired primary amine as a white foam in quantitative yield. ¹H-NMR (CDCl₃): 7.60 (d, 2H), 7.39-7.18 (m, 5H), 6.73 (d, 2H), 5.70 (d, 1H), 5.30 (d, 1H), 5.00 (q, 1H), 4.27 (s, 2H), 4.18-3.65 (m, 5H), 3.56 (dd, 1H), 3.28-2.20 (m, 12H), 2.00-1.20 (m, 6H), 0.89 (s, 6H). MS(ESI): 619(M+H).

10

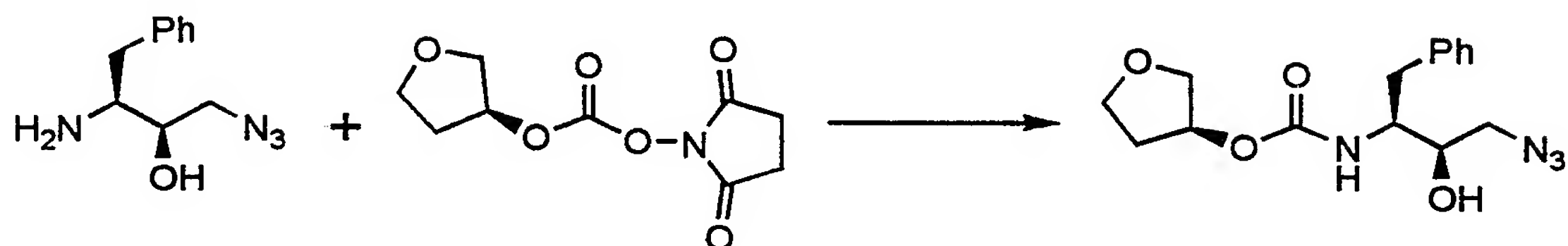
Example (Compound 293)

(3S, 3aR, 6aS) hexahydrofuro[2,3-b]furan-3-yl N-(1S, 2R)-3-
 15 [(6-amino-2,2-dimethylhexyl) (1,3-benzodioxol-5-ylsulfonyl) amino]-1-benzyl-2-hydroxypropylcarbamate
 According to example 292, steps 5 and 6,
 (3S, 3aR, 6aS) hexahydrofuro[2,3-b]furan-3-yl N-(1S, 2R)-1-benzyl-3-[(5-cyano-2,2-dimethylpentyl) amino]-2-
 20 hydroxypropylcarbamate was subjected to sulfonylation
 (with 3,4-methylenedioxybenzenesulfonyl chloride instead of 4-nitrobenzenesulfonyl chloride) followed by Raney nickel reduction to afford the desired primary amine as a white foam. ¹H-NMR (DMSO-d₆): 7.41-7.12 (m, 8H), 7.08 (d, 1H), 6.20 (s, 2H), 5.56 (d, 1H), 4.81 (q, 1H), 3.89-3.66 (m, 4H), 3.65-2.39 (m, 14H), 1.90 (m, 1H), 1.71 (m, 1H), 1.50-1.12 (m, 6H), 0.93 (s, 6H). MS(ESI): 648(M+H).

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Example (Compound 295)

5 Step 1:

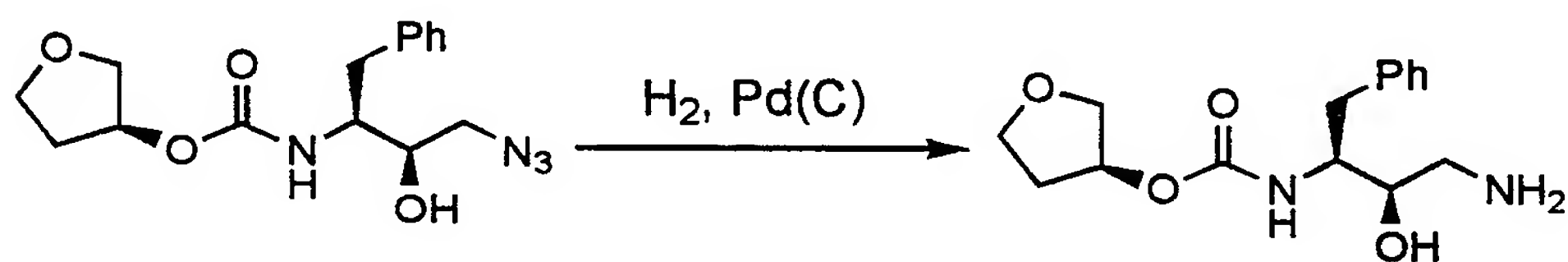


(3S)-tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-azido-2-hydroxypropyl) carbamate

10 Obtained in a similar fashion to Example 292, step 2, using the corresponding activated carbonate (see US 5,585,397). MS (ESI): 321 (M+H).

Step 2:

15

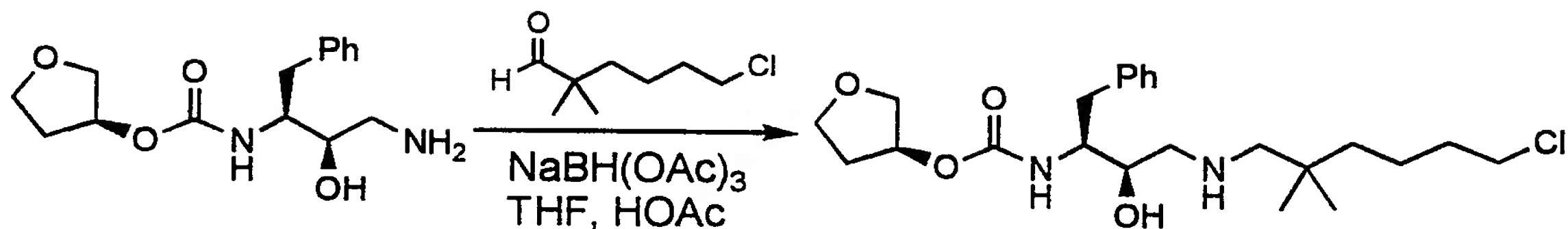


(3S)-tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-amino-2-hydroxypropyl) carbamate

20 The reduction was carried out according to example 257, step 4, to afford the desired amine as a white crystalline solid. MS (ESI): 294 (M+H).

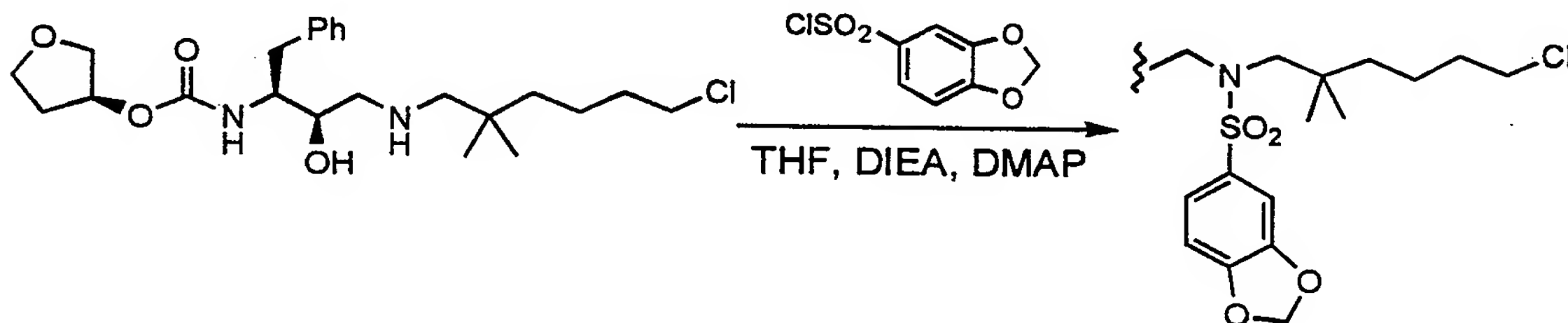
Step 3:

-294-



5 **(3S)-tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-[(6-chloro-2,2-dimethylhexyl)amino]-2-hydroxypropyl)carbamate**
 According to example 257, step 5, (3S)-tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-amino-2-hydroxypropyl)carbamate was subjected to reductive alkylation with 6-chloro-2,2-dimethylhexanal to afford the desired compound
 10 as a viscous oil in 93% yield. MS(ESI): 441(M+H).

Step 4:



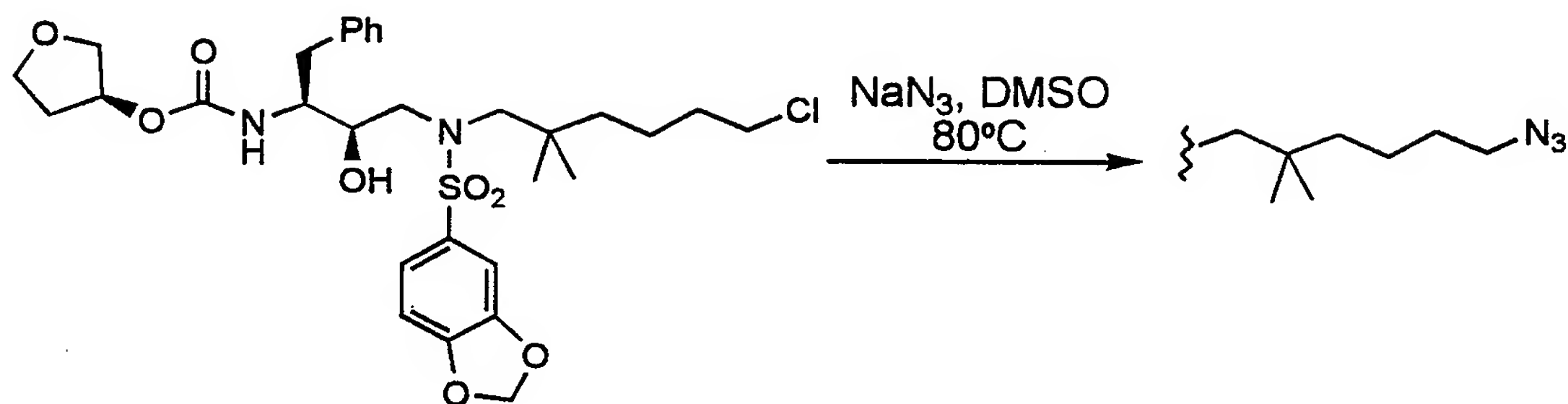
15

(3S)-tetrahydro-3-furanyl N-((1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-chloro-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropyl)carbamate
 According to example 257, step 6, (3S)-tetrahydro-3-furanyl N-((1S,2R)-1-benzyl-3-[(6-chloro-2,2-dimethylhexyl)amino]-2-hydroxypropyl)carbamate was
 20 subjected to sulfonylation to afford the desired sulfonamide as a white foam in 65% yield. MS(ESI): 625(M+H).

25

Step 5:

-295-

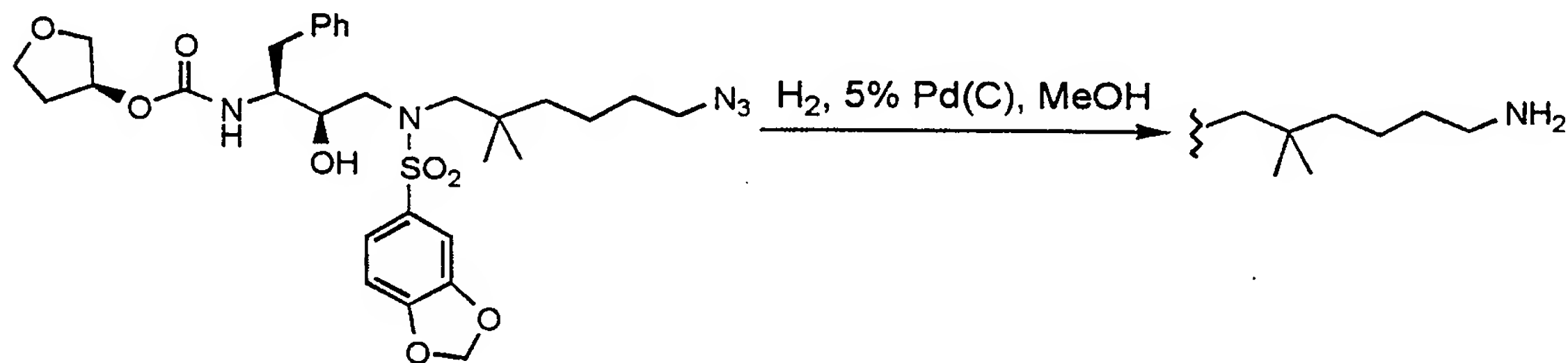


(3S)-tetrahydro-3-furanyl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-azido-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropylcarbamate

A solution of 0.320 g (0.512 mmol) of (3S)-tetrahydro-3-furanyl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-chloro-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropylcarbamate and 66 mg (1.02 mmol) of sodium azide in 4 mL of DMSO was heated to 80°C with stirring. After 2.5 hours the solution was cooled to RT and diluted with CH₂Cl₂, washed with water (3x), dried over MgSO₄, and concentrated in vacuo. The crude residue was purified by flash chromatography (SiO₂, CH₂Cl₂/MeOH) to afford 0.225 g (70%) of the desired azide as a clear viscous oil.

MS (ESI): 632 (M+H).

Step 6:

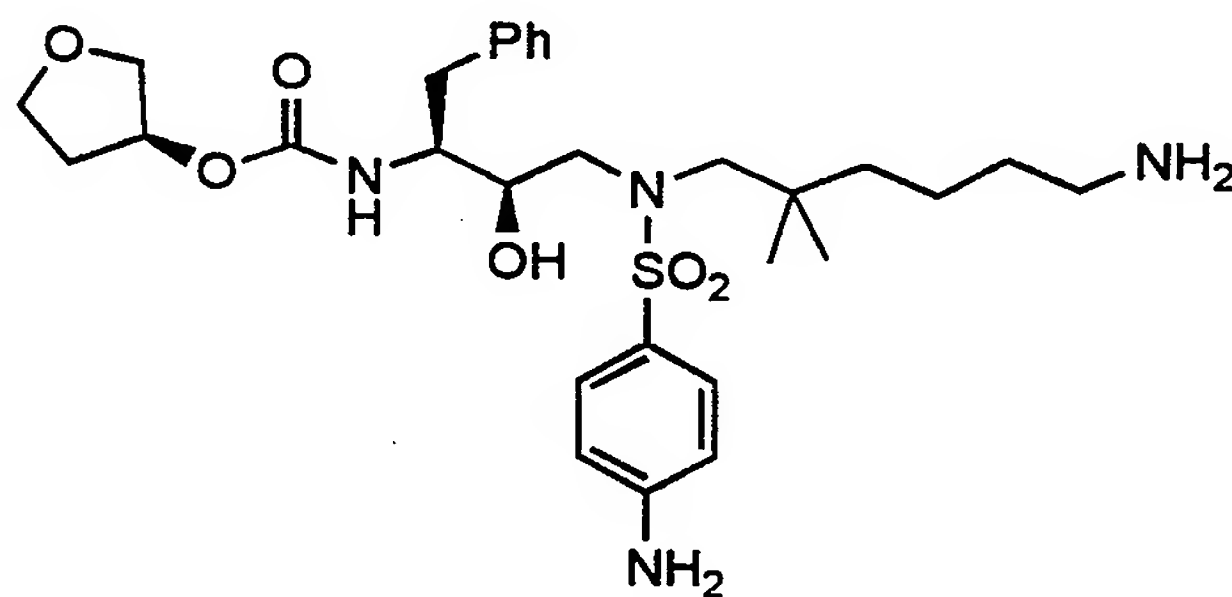


-296-

(3S)-tetrahydro-3-furanyl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-amino-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropylcarbamate

According to example 257, step 4, (3S)-tetrahydro-3-furanyl N-(1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(6-azido-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropylcarbamate was subjected to catalytic hydrogenation followed by flash chromatography (SiO₂, CH₂Cl₂/2M NH₃ in MeOH) to afford the desired amine as a white foam in 45% yield. H¹-NMR (CDCl₃): 7.32 (dd, 1H), 7.28-7.15 (m, 6H), 6.87 (d, 1H), 6.07 (s, 2H), 5.18-5.03 (m, 2H), 3.95 (t, 1H), 3.85-3.67 (m, 4H), 3.54 (d, 1H), 3.20-2.30 (m, 11H), 2.08 (m, 1H), 1.88 (m, 1H), 1.48-1.21 (m, 6H), 0.89 (d, 6H). MS (ESI): 606 (M+H).

Example (Compound 296)



(3S)-tetrahydro-3-furanyl N-(1S,2R)-3-[(4-aminophenylsulfonyl)(6-amino-2,2-dimethylhexyl)amino]-1-benzyl-2-hydroxypropylcarbamate

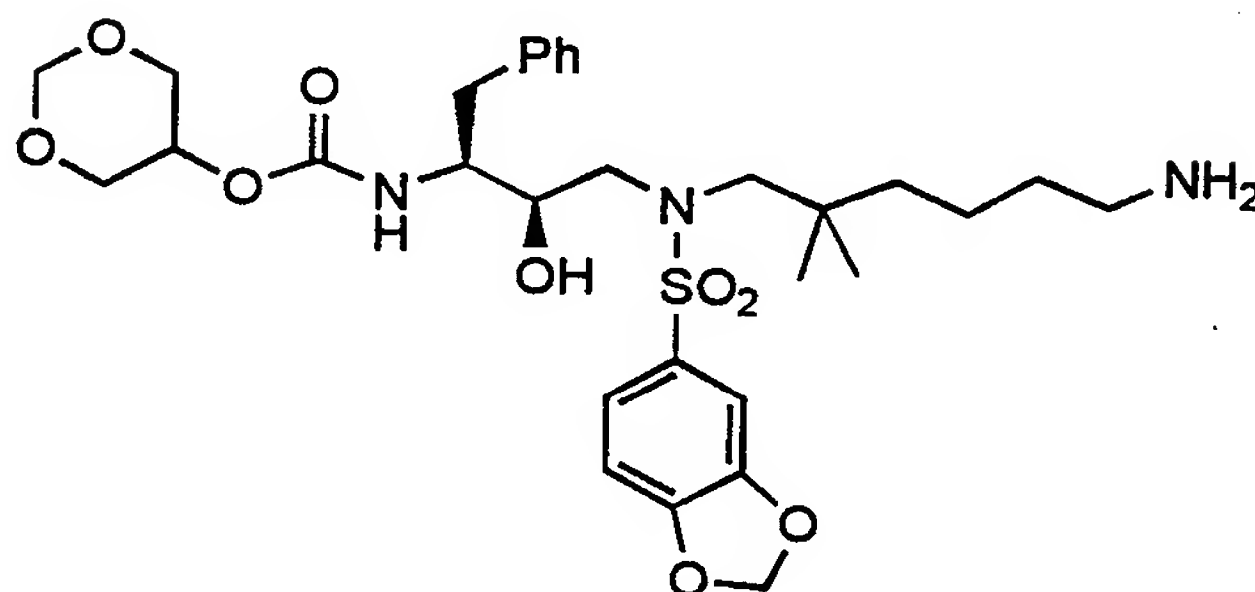
The title compound was prepared according to example 295 with the exception that 4-nitrobenzenesulfonyl chloride was used in step 4. H¹-NMR (CDCl₃): 7.53 (d, 2H), 7.30-7.13 (m, 5H), 6.66 (d, 2H), 5.13 (d, 1H), 5.06 (br s, 1H), 4.21 (s, 2H), 3.97 (t, 1H), 3.89-3.65 (m, 4H), 3.52

-297-

(d, 1H), 3.20-2.30 (m, 11H), 2.04 (m, 1H), 1.88 (m, 1H),
1.48-1.15 (m, 6H), 0.87 (d, 6H). MS(ESI): 577(M+H).

5

Example (Compound 297)

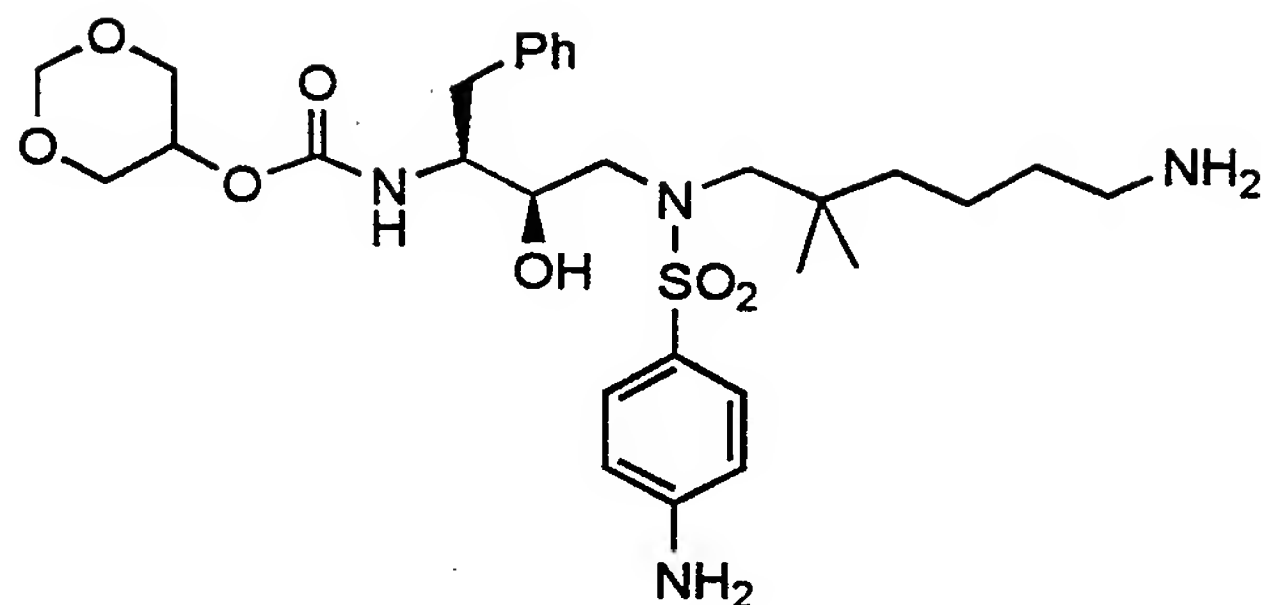


10 [1,3]-dioxan-5-yl N-(1S,2R)-3-[(1,3-benzodioxol-5-
ylsulfonyl)(6-amino-2,2-dimethylhexyl)amino]-1-benzyl-2-
hydroxypropylcarbamate

The title compound was prepared according to example 295
with the exception that 1,3-dioxan-5-yl 4-nitrophenyl
carbonate (see US 5,585,397) was used as the acylating
15 reagent in step 1. ¹H-NMR (CDCl₃): 7.32 (dd, 1H), 7.28-
7.14 (m, 6H), 6.87 (d, 1H), 6.06 (s, 2H), 5.46 (d, 1H),
4.89 (d, 1H), 4.72 (d, 1H), 4.43 (br s, 1H), 3.98 (t,
1H), 3.94-3.72 (m, 4H), 3.68 (d, 1H), 3.21-2.50 (m, 11H),
1.49-1.38 (m, 2H), 1.33-1.21 (m, 4H), 0.89 (d, 6H).
20 MS(ESI): 622(M+H).

Example (Compound 298)

-298-

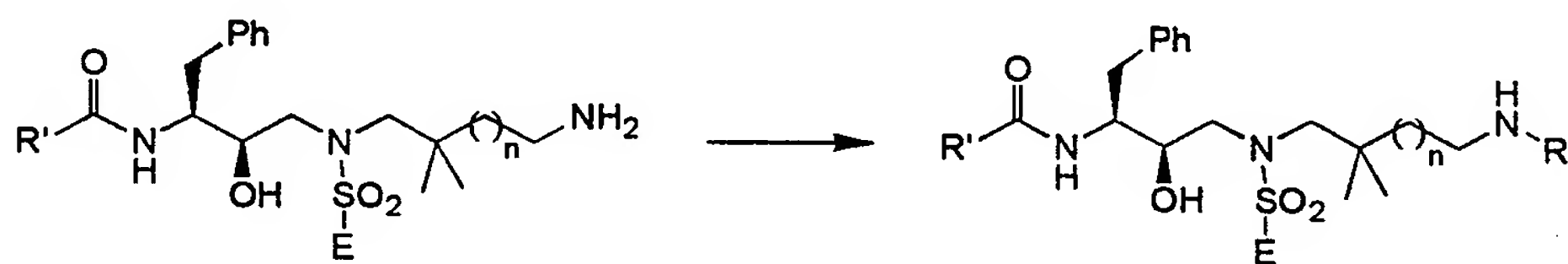


[1,3]-dioxan-5-yl N-(1S,2R)-3-[(4-aminophenylsulfonyl) (6-amino-2,2-dimethylhexyl) amino]-1-benzyl-2-

5 hydroxypropylcarbamate

The title compound was prepared according to example 295 with the exceptions that 1,3-dioxan-5-yl 4-nitrophenyl carbonate (see US 5,585,397) was used as the acylating reagent in step 1 and 4-nitrobenzenesulfonyl chloride was used as the sulfonylating reagent in step 4. H¹-NMR (CDCl₃): 7.52 (d, 2H), 7.30-7.13 (m, 5H), 6.67 (d, 2H), 5.40 (d, 1H), 4.89 (d, 1H), 4.70 (d, 1H), 4.43 (br s, 1H), 4.20 (s, 2H), 4.00 (t, 1H), 3.94-3.73 (m, 4H), 3.66 (d, 1H), 3.12 (dd, 1H), 3.07-2.40 (m, 10H), 1.45-1.33 (m, 2H), 1.30-1.19 (m, 6H), 0.88 (d, 6H). MS(ESI): 593(M+H).

Examples (Compounds 299-372)



20 General procedure for reactions of primary amine scaffolds with various electrophiles:

A solution of the primary amine (0.02 M) in anhydrous THF at 0°C was treated with 1.05 equiv. of N,N-diisopropylethylamine (omitted for reactions with

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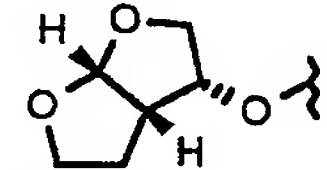
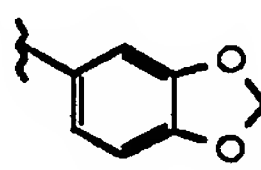
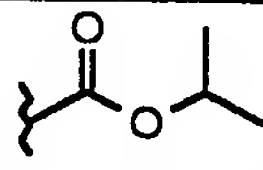
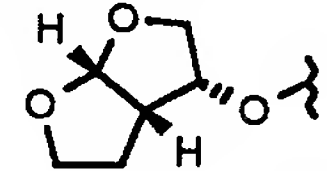
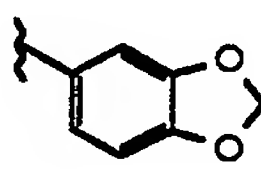
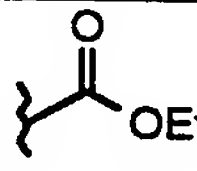
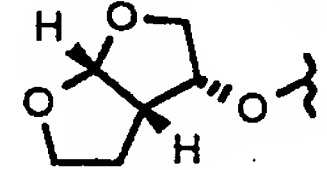
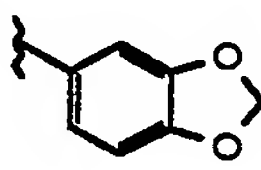
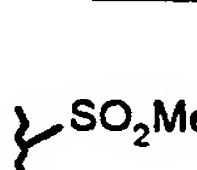
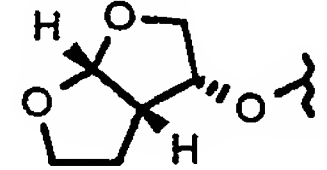
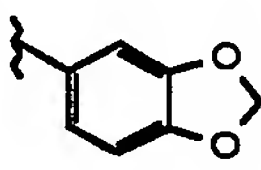
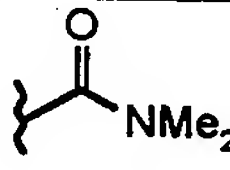
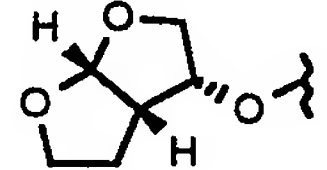
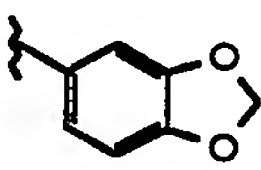
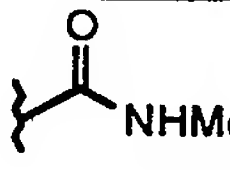
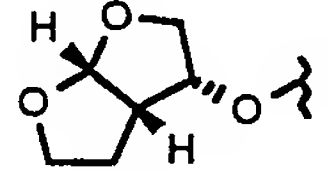
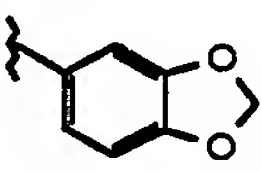
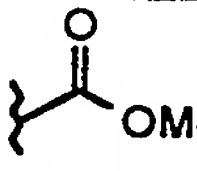
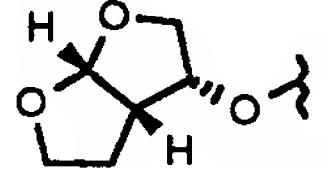
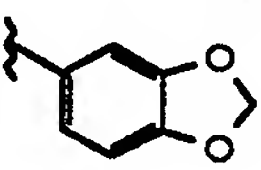
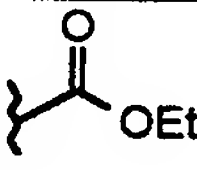
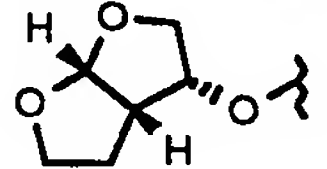
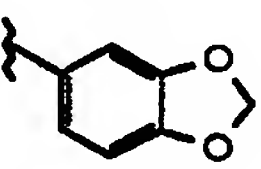
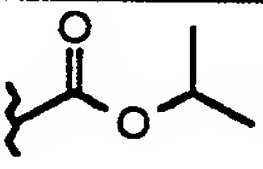
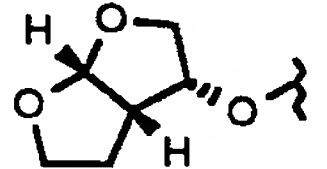
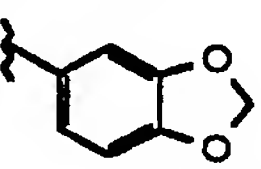
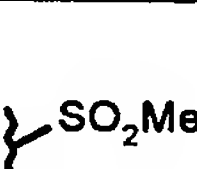
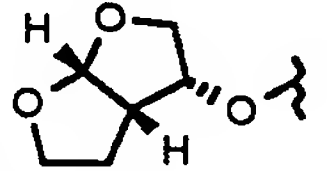
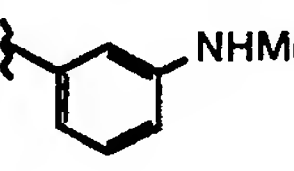
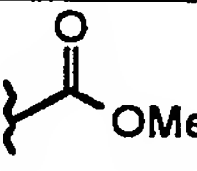
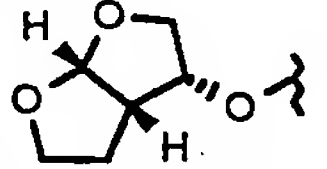
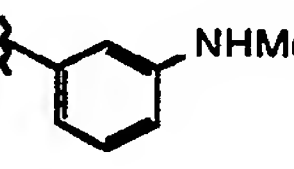
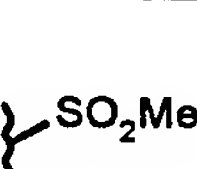
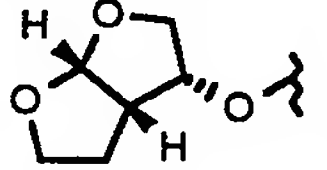
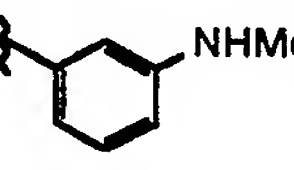
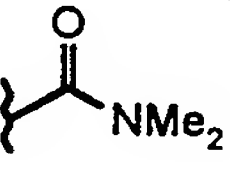
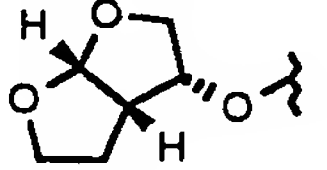
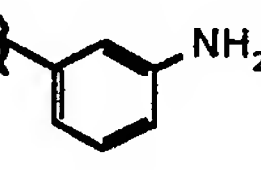
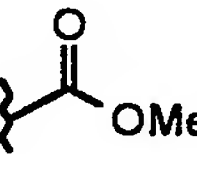
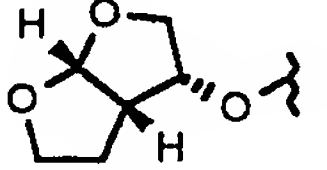
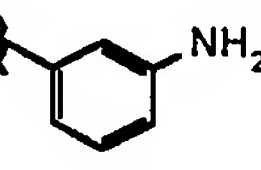
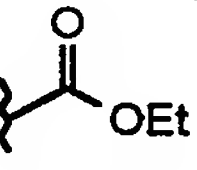
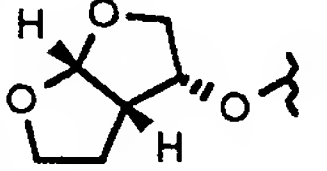
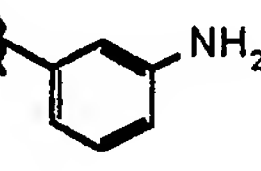
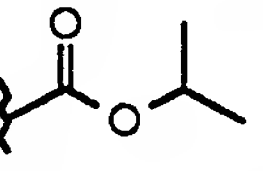
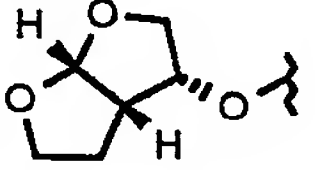
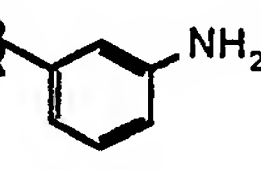
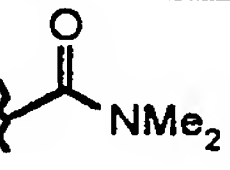
isocyanates, examples 301, 306, and 312) followed by 1.05 equiv. of the appropriate electrophile (chloroformate, sulfonyl chloride, carbamyl chloride, or isocyanate). The resulting solution was allowed to warm to RT with stirring. When analysis by TLC indicated the reaction to be complete the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, CH₂Cl₂/MeOH or CH₂Cl₂/2M NH₃ in MeOH) to afford the purified product.

10

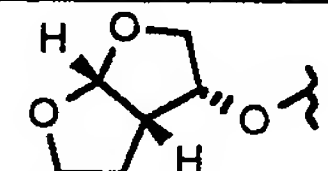
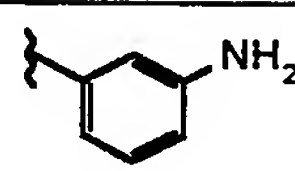
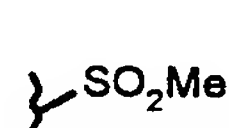
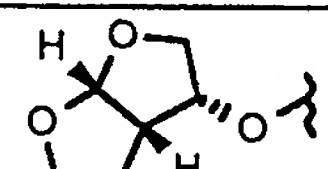
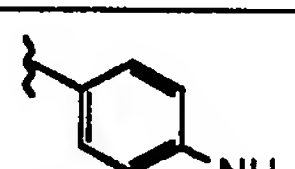
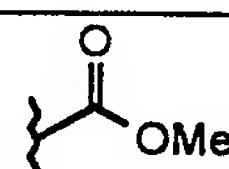
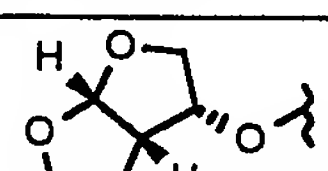
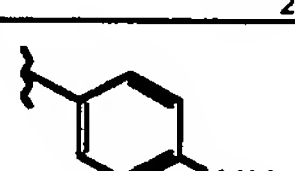
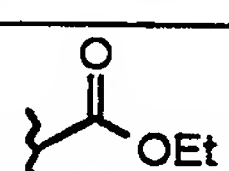
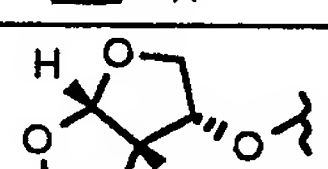
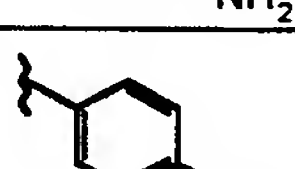

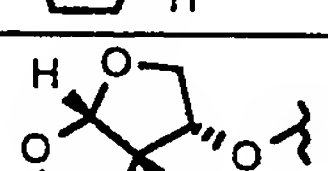
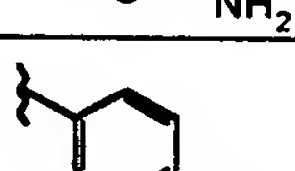
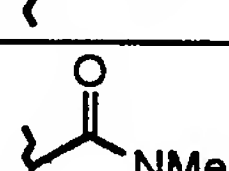
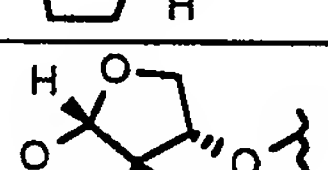
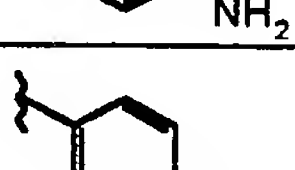
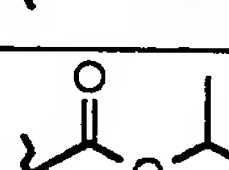
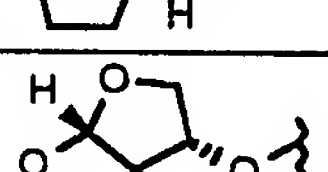
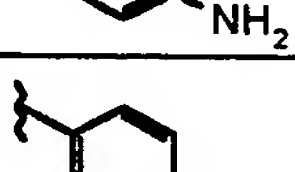
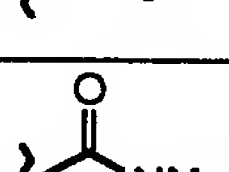
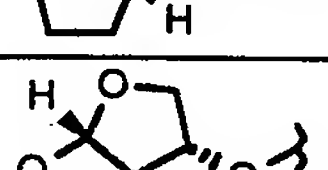
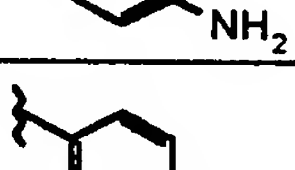
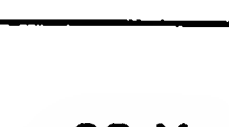
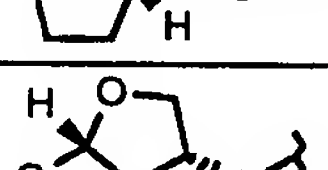
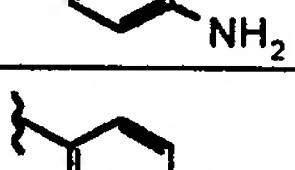
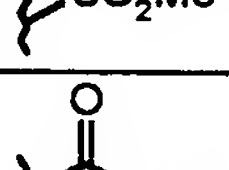
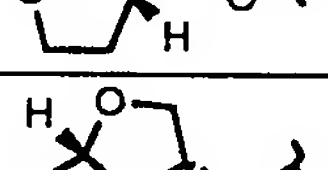
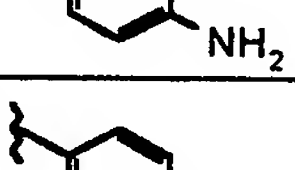
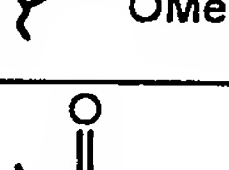
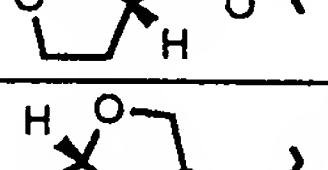
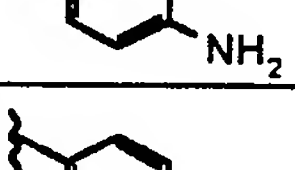
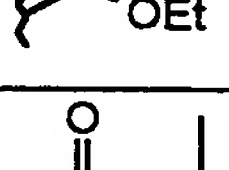
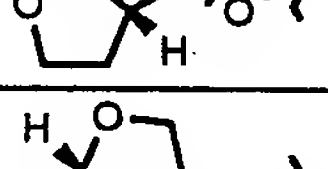
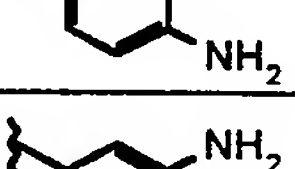
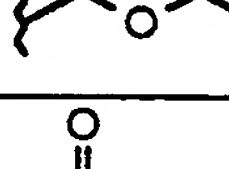
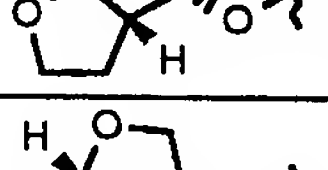
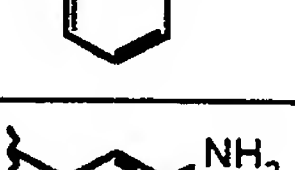
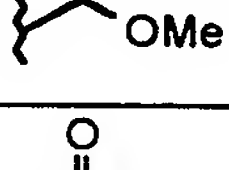
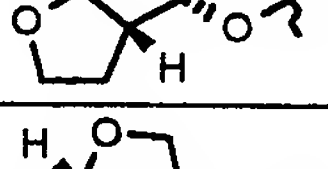

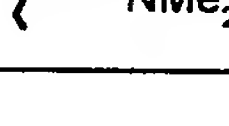
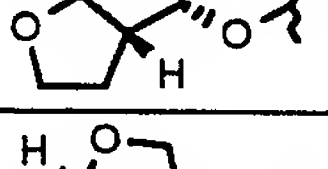

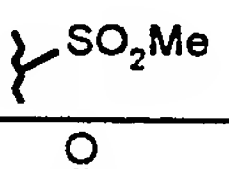
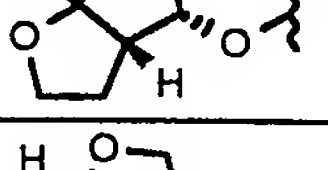
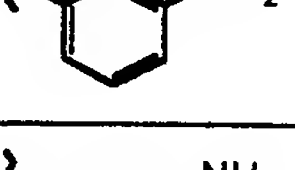
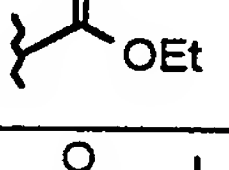
Mass Spectral Data for Compounds 299-372:

Example	Amine Precursor	R'	E	n	R	MS (ESI)
299	244			3		720 (M+H)
300	244			3		734 (M+H)
301	244			3		727 (M+Na)
302	257			2		706 (M+H)
303	257			2		720 (M+H)
304	257			2		730 (M+Na)
305	257			2		744 (M+Na)
306	257			2		713 (M+Na)
307	282			1		678 (M+H)

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308	282			1		706 (M+H)
309	282			1		714 (M+Na)
310	282			1		720 (M+Na)
311	282			1		691 (M+H)
312	282			1		699 (M+Na)
313	283			4		720 (M+H)
314	283			4		734 (M+H)
315	283			4		748 (M+H)
316	283			4		762 (M+Na)
317	284			1		663 (M+H)
318	284			1		705 (M+Na)
319	284			1		676 (M+H)
320	285			1		649 (M+H)
321	285			1		663 (M+H)
322	285			1		677 (M+H)
323	285			1		662 (M+H)

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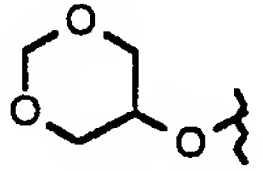
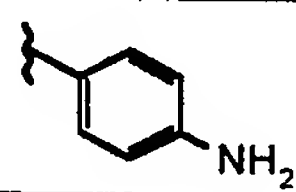
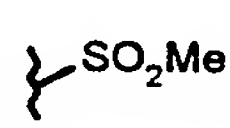
324	285			1		669 (M+H)
325	286			1		649 (M+H)
326	286			1		663 (M+H)
327	286			1		691 (M+H)
328	286			1		662 (M+H)
329	286			1		677 (M+H)
330	287			2		676 (M+H)
331	287			2		705 (M+Na)
332	287			2		663 (M+H)
333	287			2		699 (M+Na)
334	287			2		691 (M+H)
335	288			2		685 (M+Na)
336	288			2		698 (M+Na)
337	288			2		705 (M+Na)
338	288			2		699 (M+Na)
339	288			2		691 (M+H)

340	289			3		711 (M+H)
341	289			3		705 (M+H)
342	289			3		719 (M+H)
343	290			3		677 (M+H)
344	290			3		691 (M+H)
345	290			3		705 (M+H)
346	290			3		690 (M+H)
347	290			3		697 (M+H)
348	291			3		677 (M+H)
349	291			3		691 (M+H)
350	291			3		705 (M+H)
351	291			3		690 (M+H)
352	291			3		697 (M+H)
353	291			3		693 (M+H)
354	291			3		707 (M+H)
355	292			3		677 (M+H)

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356	292			3		691 (M+H)
357	292			3		705 (M+H)
358	292			3		697 (M+H)
359	293			3		728 (M+Na)
360	293			3		720 (M+H)
361	293			3		734 (M+H)
362	293			3		726 (M+H)
363	294			3		721 (M+H)
364	294			3		741 (M+H)
365	295			3		664 (M+H)
366	295			3		684 (M+H)
367	296			3		635 (M+H)
368	296			3		655 (M+H)
369	297			3		680 (M+H)
370	297			3		722 (M+Na)
371	298			3		651 (M+H)

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372	298			3		671 (M+H)
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Proton NMR data for selected compounds from the above table:

5

Example (Compound 299) NMR (CDCl₃): 7.32 (dd, 1H), 7.29-7.14 (m, 6H), 6.89 (d, 1H), 6.07 (s, 2H), 5.60 (d, 1H), 5.24 (m, 1H), 4.98 (q, 1H), 4.81 (br s, 1H), 4.09 (q, 2H), 4.01 (m, 1H), 3.91 (dd, 1H), 3.80 (m, 2H), 3.63 (m, 2H), 3.21-3.01 (m, 5H), 2.96 (d, 1H), 2.90-2.65 (m, 3H), 1.70-1.17 (m, 11H), 0.89 (d, 6H).

Example (Compound 300) NMR (CDCl₃): 7.32 (dd, 1H), 7.28-7.11 (m, 6H), 6.89 (d, 1H), 6.08 (s, 2H), 5.60 (d, 1H), 5.29 (br s, 1H), 4.97 (q, 1H), 4.89 (m, 1H), 4.72 (br s, 1H), 4.01 (t, 1H), 3.91 (dd, 1H), 3.79 (m, 2H), 3.63 (m, 2H), 3.21-3.02 (m, 5H), 2.96 (d, 1H), 2.84 (m, 1H), 2.73 (m, 2H), 1.60-1.15 (m, 14H), 0.88 (d, 6H).

Example (Compound 307) NMR (CDCl₃): 7.32 (dd, 1H), 7.28-7.13 (m, 6H), 6.89 (d, 1H), 6.08 (s, 2H), 5.60 (d, 1H), 5.25 (m, 1H), 5.08 (br s, 1H), 4.96 (q, 1H), 4.20-3.95 (m, 2H), 3.91 (dd, 1H), 3.81 (m, 2H), 3.64 (m, 5H), 3.25-2.92 (m, 6H), 2.87 (m, 1H), 2.81-2.66 (m, 2H), 1.70-1.30 (m, 4H), 0.98 (s, 3H), 0.91 (s, 3H).

Example (Compound 309) NMR (CDCl₃): 7.32 (dd, 1H), 7.28-7.10 (m, 6H), 6.88 (d, 1H), 6.08 (s, 2H), 5.60 (d, 1H), 5.25 (m, 1H), 5.00 (m, 2H), 4.20-3.89 (m, 5H), 3.80 (m, 2H), 3.63 (m, 2H), 3.28-2.93 (m, 6H), 2.88 (m, 1H), 2.82-

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2.65 (m, 2H), 1.80-1.30 (m, 4H), 1.21 (t, 3H), 0.99 (s, 3H), 0.91 (s, 3H).

5 Example (Compound 313) NMR (CDCl₃): 7.32 (dd, 1H), 7.30-7.12 (m, 6H), 6.89 (d, 1H), 6.09 (s, 2H), 5.61 (d, 1H), 4.97 (m, 2H), 4.83 (br s, 1H), 4.03 (m, 1H), 3.92 (m, 1H), 3.82 (m, 2H), 3.64 (m, 5H), 3.21-2.67 (m, 9H), 1.70-1.33 (m, 4H), 1.23 (m, 6H), 0.87 (d, 6H).

10 Example (Compound 314) NMR (CDCl₃): 7.33 (dd, 1H), 7.29-7.13 (m, 6H), 6.89 (d, 1H), 6.09 (s, 2H), 5.60 (d, 1H), 4.98 (m, 2H), 4.76 (br s, 1H), 4.13-4.00 (m, 3H), 3.91 (dd, 1H), 3.82 (m, 2H), 3.66 (m, 2H), 3.21-2.68 (m, 9H), 1.66-1.33 (m, 5H), 1.23 (m, 8H), 0.89 (s, 6H).

15

Example (Compound 315) NMR (CDCl₃): 7.32 (dd, 1H), 7.28-7.13 (m, 6H), 6.90 (d, 1H), 6.08 (s, 2H), 5.61 (d, 1H), 4.98 (m, 2H), 4.88 (m, 1H), 4.67 (br s, 1H), 4.03 (m, 1H), 3.91 (dd, 1H), 3.82 (m, 2H), 3.66 (m, 2H), 3.21-2.66 (m, 9H), 1.70-1.31 (m, 6H), 1.24 (m, 10H), 0.89 (s, 6H).

20

Example (Compound 316) NMR (CDCl₃): 7.32 (dd, 1H), 7.30-7.12 (m, 6H), 6.90 (d, 1H), 6.09 (s, 2H), 5.61 (d, 1H), 4.98 (m, 2H), 4.62 (m, 1H), 4.04 (m, 2H), 4.95-3.76 (m, 3H), 3.68 (m, 2H), 3.21-3.05 (m, 4H), 3.04-2.82 (m, 6H), 2.73 (m, 2H), 1.56 (m, 3H), 1.43-1.20 (m, 7H), 0.90 (d, 6H).

25

Example (Compound 359) NMR (CDCl₃): 3.36-7.13 (m, 7H), 6.90 (d, 1H), 6.09 (s, 2H), 5.64 (d, 1H), 5.11 (d, 1H), 4.97 (q, 1H), 4.87 (br s, 1H), 4.07 (br s, 2H), 3.91 (m, 2H), 3.79 (m, 2H), 3.66 (s, 3H), 3.50 (m, 1H), 3.22-2.70

30

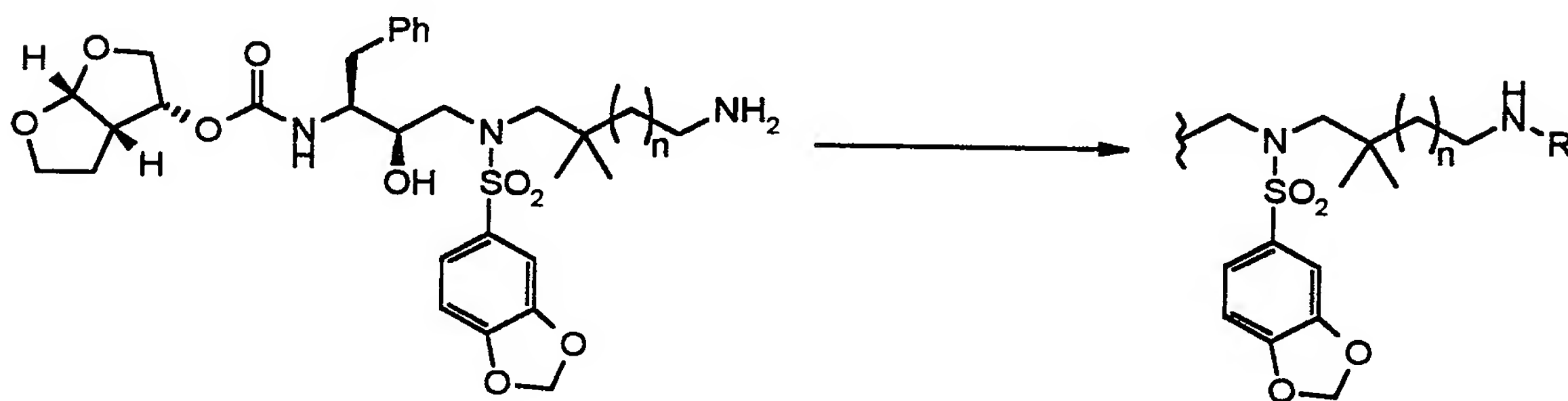
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(m, 9H), 1.94 (m, 1H), 1.80 (m, 1H), 1.47 (br s, 2H),
1.24 (br s, 4H), 0.89 (d, 6H).

Example (Compound 360) NMR (CDCl₃): 7.36-7.12 (m, 7H),
5 6.88 (d, 1H), 6.09 (s, 2H), 5.64 (d, 1H), 5.14 (d, 1H),
4.96 (q, 1H), 4.80 (br s, 1H), 4.09 (br s, 4H), 3.91 (m,
2H), 3.76 (m, 2H), 3.49 (m, 1H), 3.21-2.68 (m, 9H), 1.95
(m, 1H), 1.80 (m, 1H), 1.47 (br s, 2H), 1.23 (m, 7H),
0.89 (d, 6H).

10

Examples (Compounds 373-519)



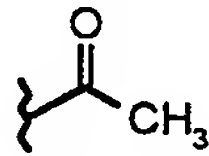
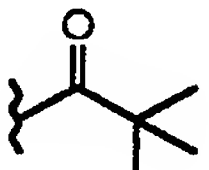
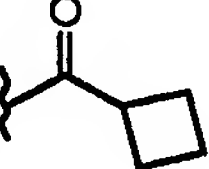
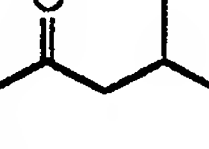
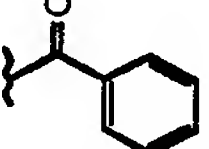
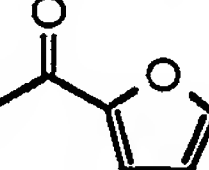
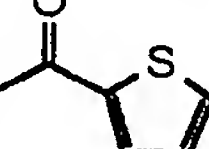
15 *General procedure for reactions of primary amine
scaffolds with various electrophiles using scavenger
resins:*

The following reactions were carried out in parallel, in
sets ranging from 9 to 48, and were done in teflon-capped
20 glass vials using a rotary shaker for agitation. To
vials containing 5 to 10 equiv. of either Amberlyst A-21
resin (Aldrich, prepared by washing twice each with MeOH,
CH₂Cl₂, THF, and then drying in vacuo at 60°C overnight)
or PS-DIEA resin (Argonaut Technologies, prepared as
25 above) were added 2 mL of 16 mM solutions of the primary
amine starting materials in anhydrous THF or anhydrous
CH₂Cl₂. These resins were not employed when isocyanates

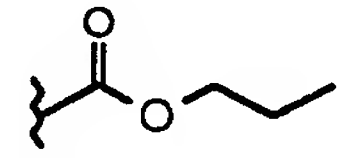
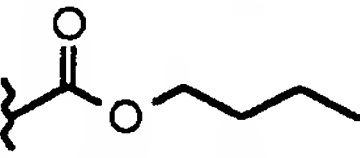
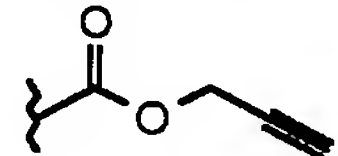
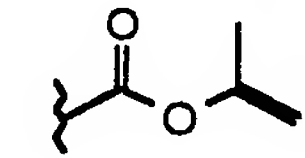
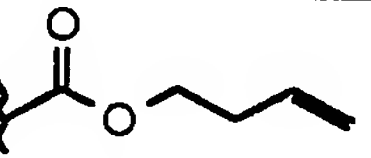
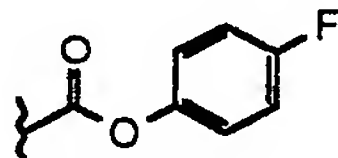
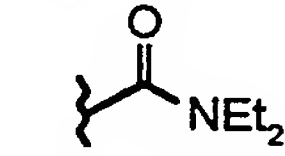
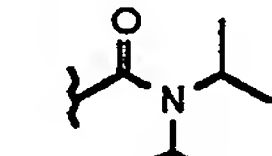
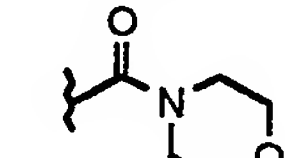
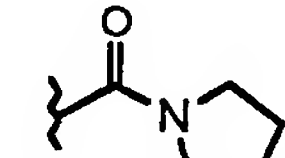
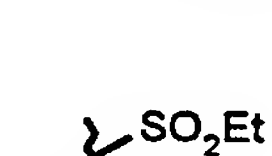
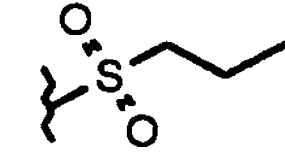
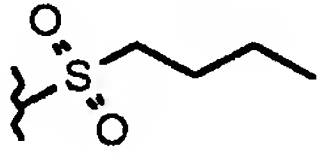
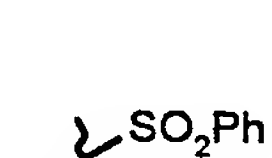
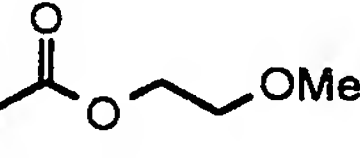
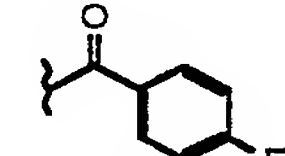
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or isothiocyanates were used as electrophiles. This was followed by addition of 3 to 5 equivalents of electrophile (acid chlorides, chloroformates, carbamyl chlorides, sulfonyl chlorides, isocyanates, isothiocyanates) either neat or as solutions in anhydrous CH_2Cl_2 . The resulting mixtures were shaken at RT for 18 hours. The mixtures were then diluted two-fold with CH_2Cl_2 and treated with 5-10 equiv. of PS-Trisamine resin (Argonaut Technologies, prepared as above). This was followed by shaking at RT for an additional 18 hours. The resins were then removed by filtration and the filtrates concentrated to dryness under a stream of nitrogen to afford the desired compounds.

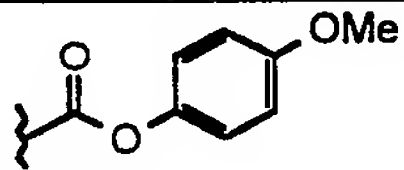
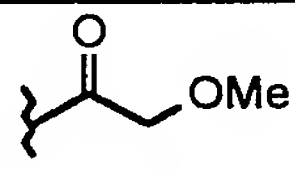
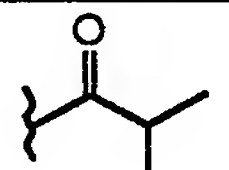
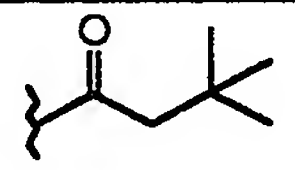
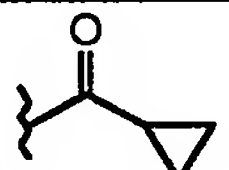
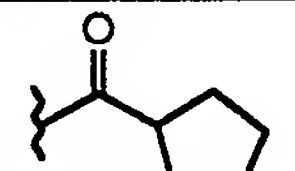
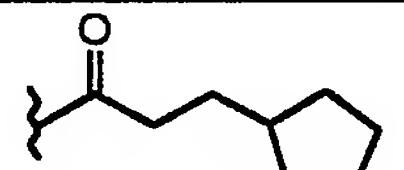
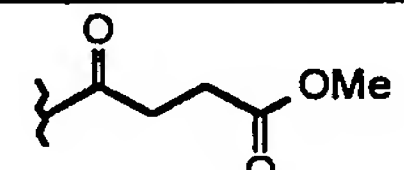
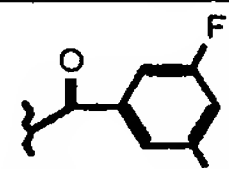
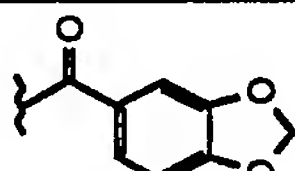
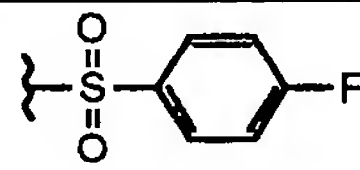
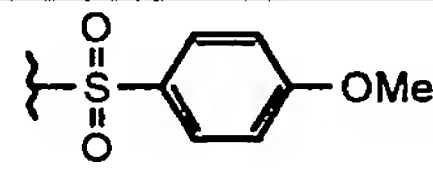
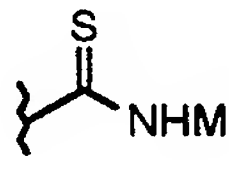
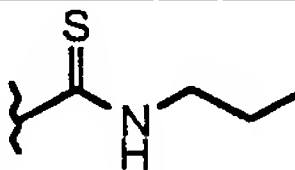
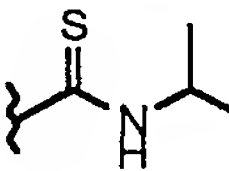
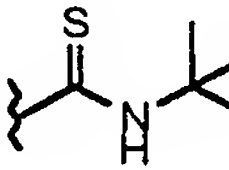
15 Mass Spectral Data for Compounds 373-519:

Example	Amine Precursor	n	R	MS (ESI)
373	282	1		683 (M+Na)
374	282	1		726 (M+Na)
375	282	1		724 (M+Na)
376	282	1		726 (M+Na)
377	282	1		746 (M+Na)
378	282	1		736 (M+Na)
379	282	1		752 (M+Na)

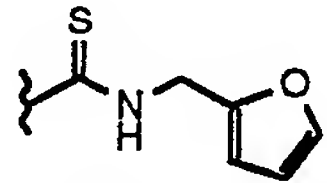
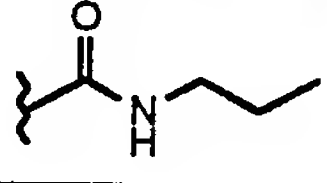
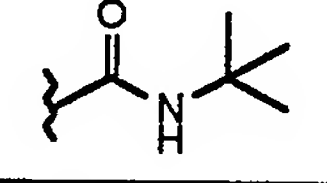
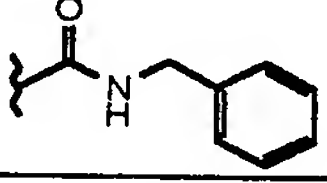
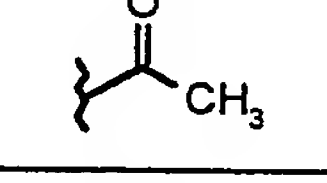
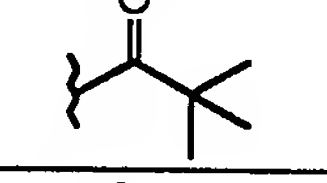
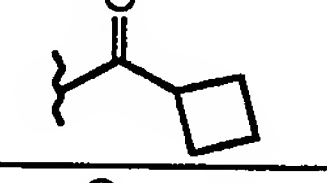
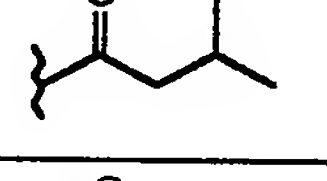
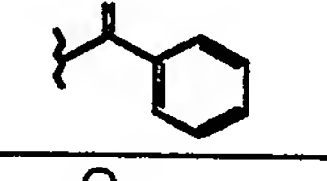
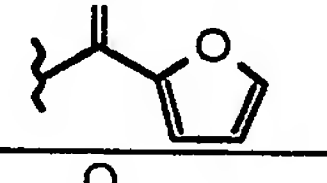
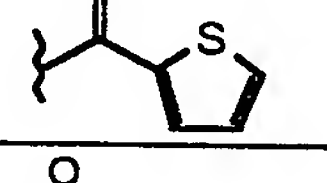
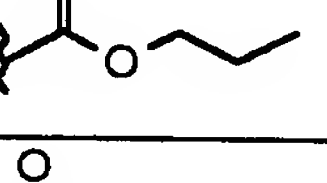
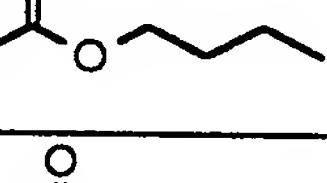
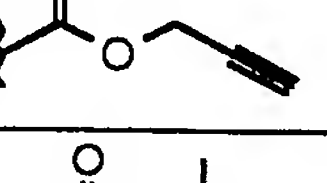
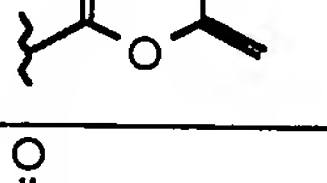
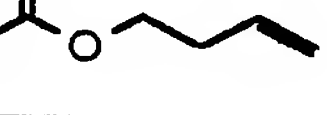
-308-

380	282	1		706 (M+H)
381	282	1		720 (M+H)
382	282	1		702 (M+H)
383	282	1		704 (M+H)
384	282	1		718 (M+H)
385	282	1		758 (M+H)
386	282	1		719 (M+H)
387	282	1		747 (M+H)
388	282	1		732 (M+H)
389	282	1		717 (M+H)
390	282	1		712 (M+H)
391	282	1		726 (M+H)
392	282	1		740 (M+H)
393	282	1		760 (M+H)
394	282	1		722 (M+H)
395	282	1		742 (M+H)

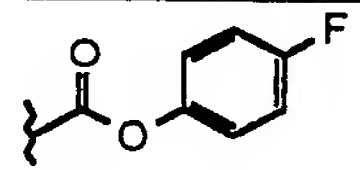
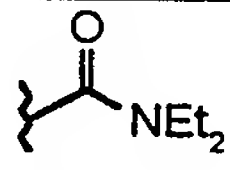
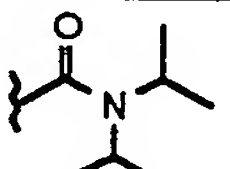
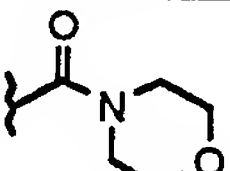
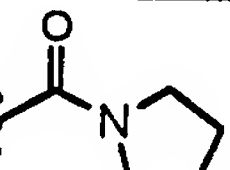
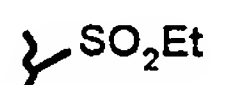
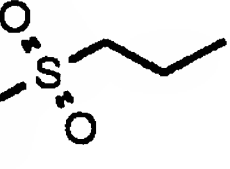
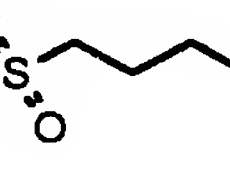
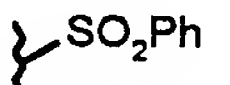
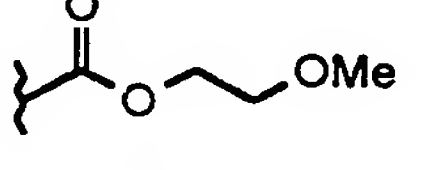
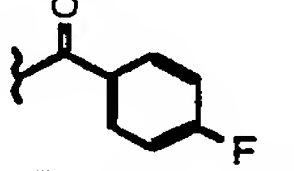
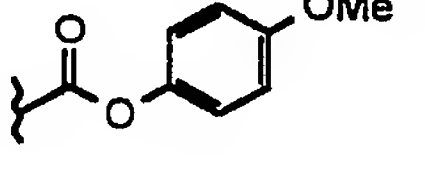
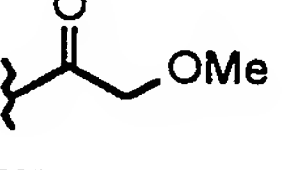
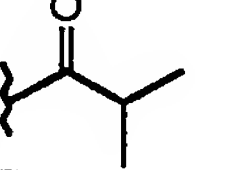
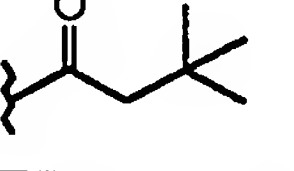
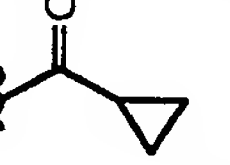
-309-

396	282	1		770 (M+H)
397	282	1		692 (M+H)
398	282	1		690 (M+H)
399	282	1		718 (M+H)
400	282	1		688 (M+H)
401	282	1		716 (M+H)
402	282	1		744 (M+H)
403	282	1		734 (M+H)
404	282	1		760 (M+H)
405	282	1		768 (M+H)
406	282	1		778 (M+H)
407	282	1		790 (M+H)
408	282	1		693 (M+H)
409	282	1		721 (M+H)
410	282	1		721 (M+H)
411	282	1		735 (M+H)

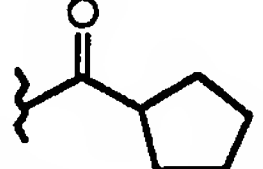
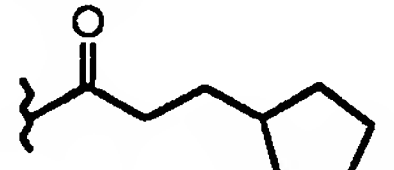
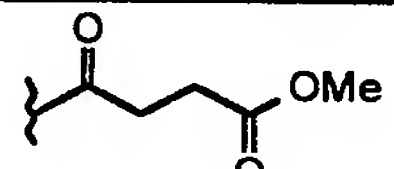
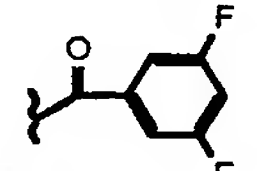
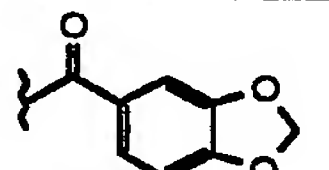
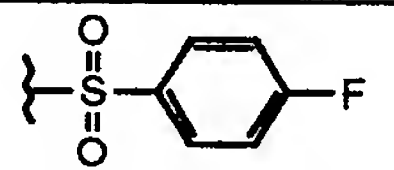
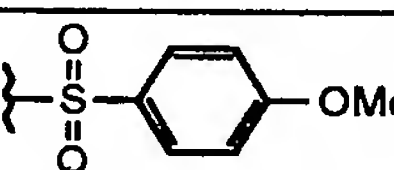
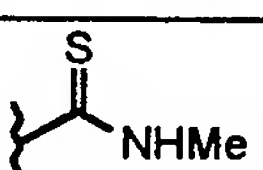
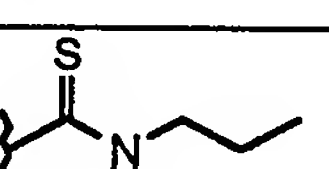
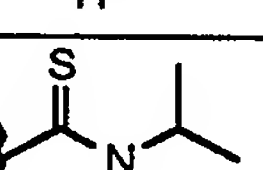
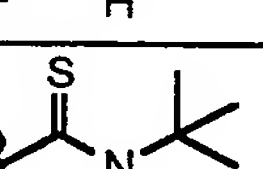
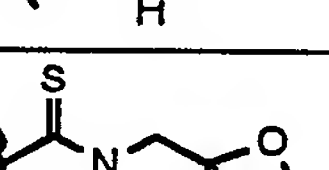
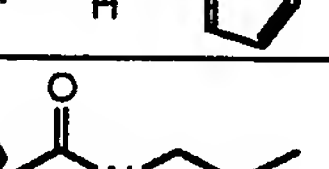
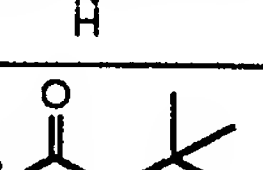
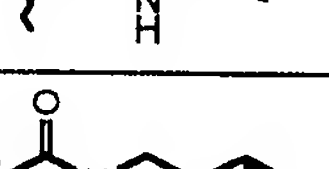
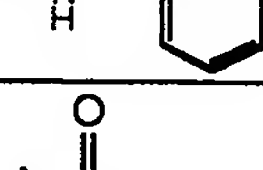
-310-

412	282	1		759 (M+H)
413	282	1		705 (M+H)
414	282	1		719 (M+H)
415	282	1		753 (M+H)
416	257	2		698 (M+Na)
417	257	2		740 (M+Na)
418	257	2		738 (M+Na)
419	257	2		740 (M+Na)
420	257	2		760 (M+Na)
421	257	2		750 (M+Na)
422	257	2		744 (M+H)
423	257	2		720 (M+H)
424	257	2		734 (M+H)
425	257	2		716 (M+H)
426	257	2		718 (M+H)
427	257	2		732 (M+H)

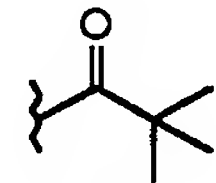
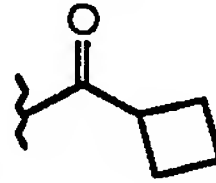
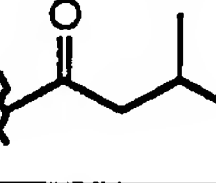
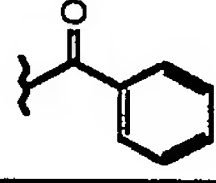
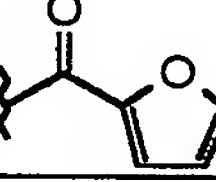
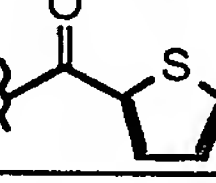
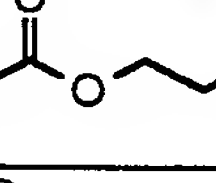
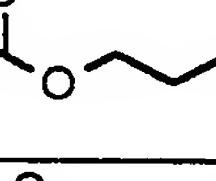
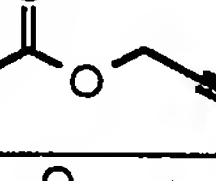
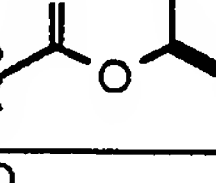
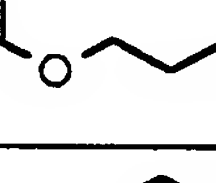
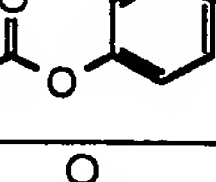
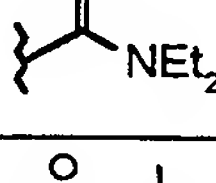
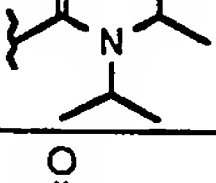
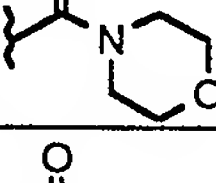
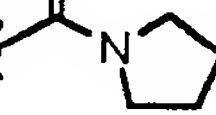
-311-

428	257	2		772 (M+H)
429	257	2		733 (M+H)
430	257	2		761 (M+H)
431	257	2		747 (M+H)
432	257	2		731 (M+H)
433	257	2		726 (M+H)
434	257	2		740 (M+H)
435	257	2		754 (M+H)
436	257	2		774 (M+H)
437	257	2		736 (M+H)
438	257	2		756 (M+H)
439	257	2		784 (M+H)
440	257	2		706 (M+H)
441	257	2		704 (M+H)
442	257	2		732 (M+H)
443	257	2		702 (M+H)


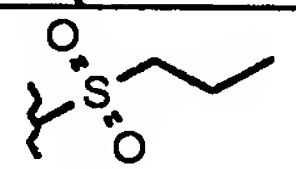
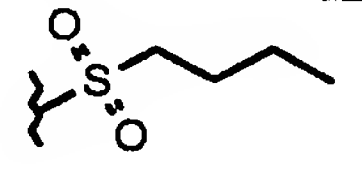
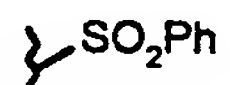
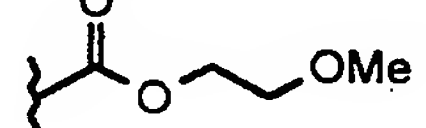
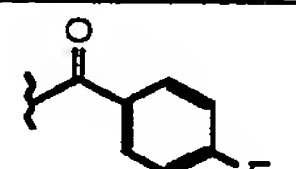
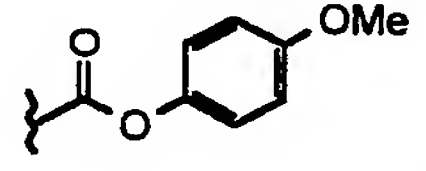
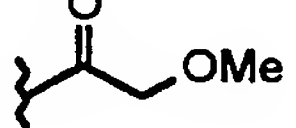
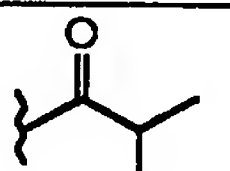
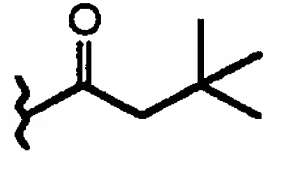
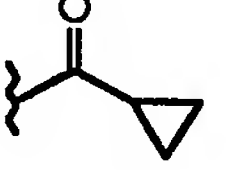
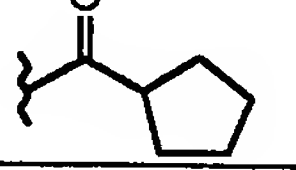
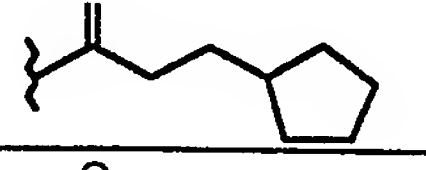
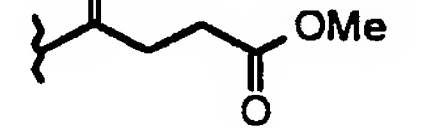
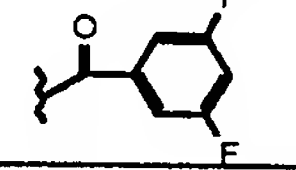
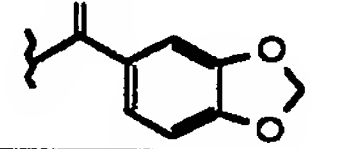
-312-

444	257	2		730 (M+H)
445	257	2		758 (M+H)
446	257	2		748 (M+H)
447	257	2		774 (M+H)
448	257	2		782 (M+H)
449	257	2		792 (M+H)
450	257	2		804 (M+H)
451	257	2		707 (M+H)
452	257	2		735 (M+H)
453	257	2		735 (M+H)
454	257	2		749 (M+H)
455	257	2		773 (M+H)
456	257	2		719 (M+H)
457	257	2		733 (M+H)
458	257	2		767 (M+H)
459	244	3		712 (M+Na)

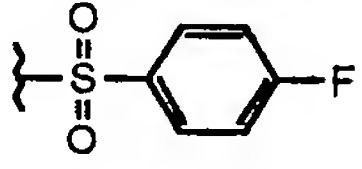
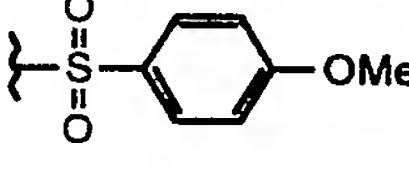
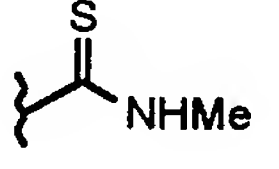
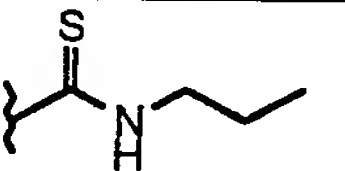
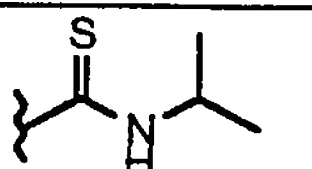
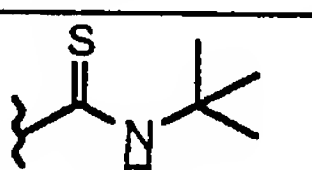
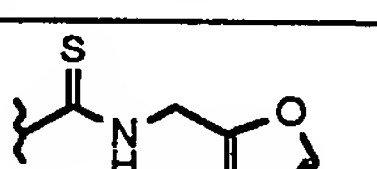
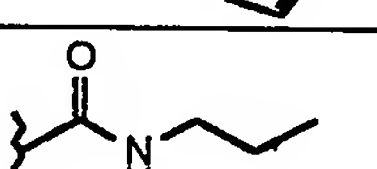
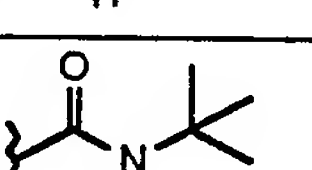
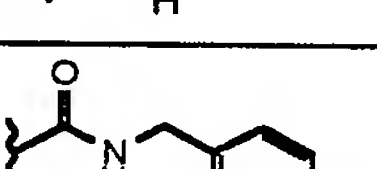
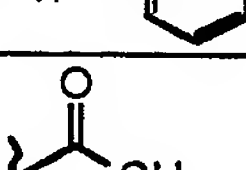
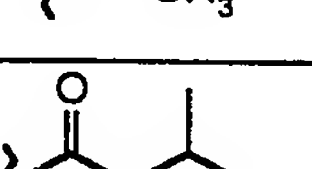
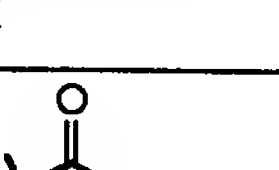
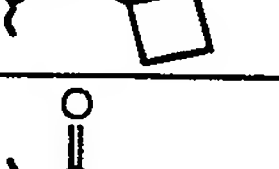
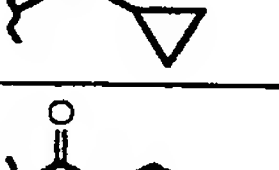
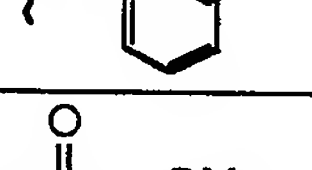
-313-

460	244	3		754 (M+Na)
461	244	3		752 (M+Na)
462	244	3		754 (M+Na)
463	244	3		774 (M+Na)
464	244	3		764 (M+Na)
465	244	3		780 (M+Na)
466	244	3		734 (M+H)
467	244	3		748 (M+H)
468	244	3		730 (M+H)
469	244	3		732 (M+H)
470	244	3		746 (M+H)
471	244	3		786 (M+H)
472	244	3		747 (M+H)
473	244	3		775 (M+H)
474	244	3		761 (M+H)
475	244	3		745 (M+H)

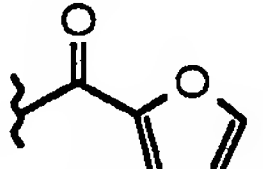
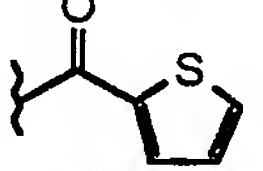
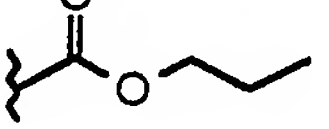
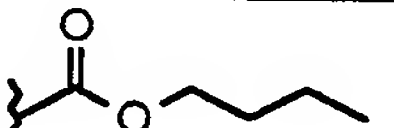
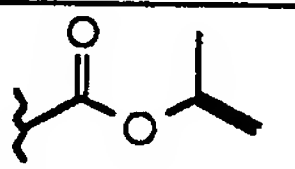
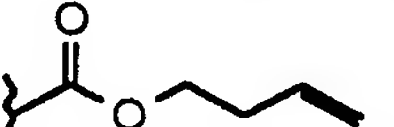
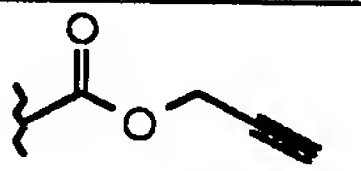
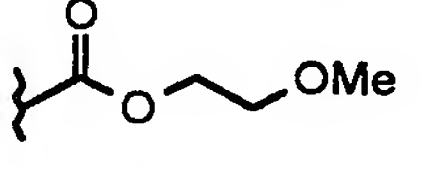
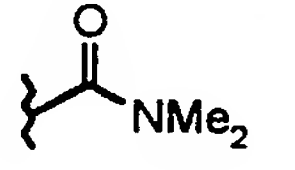
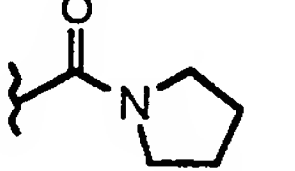
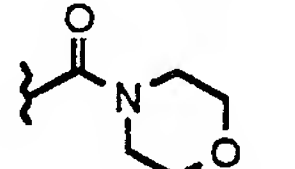
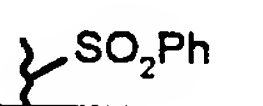
-314-

476	244	3		740 (M+H)
477	244	3		754 (M+H)
478	244	3		768 (M+H)
479	244	3		788 (M+H)
480	244	3		750 (M+H)
481	244	3		770 (M+H)
482	244	3		798 (M+H)
483	244	3		720 (M+H)
484	244	3		718 (M+H)
485	244	3		746 (M+H)
486	244	3		716 (M+H)
487	244	3		744 (M+H)
488	244	3		772 (M+H)
489	244	3		762 (M+H)
490	244	3		788 (M+H)
491	244	3		796 (M+H)

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492	244	3		806 (M+H)
493	244	3		718 (M+H)
494	244	3		721 (M+H)
495	244	3		749 (M+H)
496	244	3		749 (M+H)
497	244	3		763 (M+H)
498	244	3		787 (M+H)
499	244	3		733 (M+H)
500	244	3		747 (M+H)
501	244	3		781 (M+H)
502	283	4		704 (M+H)
503	283	4		746 (M+H)
504	283	4		744 (M+H)
505	283	4		730 (M+H)
506	283	4		766 (M+H)
507	283	4		734 (M+H)

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508	283	4		756 (M+H)
509	283	4		772 (M+H)
510	283	4		748 (M+H)
511	283	4		762 (M+H)
512	283	4		746 (M+H)
513	283	4		760 (M+H)
514	283	4		744 (M+H)
515	283	4		764 (M+H)
516	283	4		733 (M+H)
517	283	4		759 (M+H)
518	283	4		775 (M+H)
519	283	4		802 (M+H)

Proton NMR data for selected compounds from the above
5 table:

Example (Compound 375) NMR (CDCl₃): 7.39 (dd, 1H), 7.34-7.16 (m, 6H), 6.92 (d, 1H), 6.13 (s, 2H), 5.98 (br s, 1H), 5.66 (d, 1H), 5.50 (d, 1H), 5.01 (q, 1H), 4.29 (br

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s, 1H), 4.10-3.62 (m, 7H), 3.50-2.70 (m, 9H), 2.41-1.80 (m, 6H), 1.70-1.25 (m, 4H), 1.05 (s, 3H), 0.98 (s, 3H).

5 Example (Compound 378) NMR (CDCl₃): 7.46 (s, 1H), 7.39 (dd, 1H), 7.25 (m, 6H), 7.11 (m, 1H), 6.93 (d, 1H), 6.72 (t, 1H), 6.52 (m, 1H), 6.12 (s, 2H), 5.63 (d, 1H), 5.50 (d, 1H), 4.98 (q, 1H), 4.22 (br s, 1H), 4.04 (m, 1H), 3.97-2.68 (m, 14H), 1.85-1.20 (m, 4H), 1.11 (s, 3H), 1.03 (s, 3H).

10

Example (Compound 384) NMR (CDCl₃): 7.30 (dd, 1H), 7.26-7.07 (m, 6H), 6.86 (d, 1H), 6.04 (s, 2H), 5.73 (m, 1H), 5.58 (d, 1H), 5.24 (br s, 1H), 5.10-4.84 (m, 4H), 4.20-3.50 (m, 9H), 3.26-2.58 (m, 9H), 2.31 (m, 2H), 1.80-1.22 (m, 4H), 0.96 (s, 3H), 0.90 (s, 3H).

15

Example (Compound 421) NMR (CDCl₃): 7.47 (s, 1H), 0.39 (dd, 1H), 0.34-7.16 (m, 6H), 7.13 (m, 1H), 6.92 (d, 1H), 6.71 (t, 1H), 6.52 (m, 1H), 6.12 (s, 2H), 5.74 (d, 1H), 5.64 (d, 1H), 5.01 (q, 1H), 4.16 (br s, 1H), 4.10-3.60 (m, 6H), 3.59-2.68 (m, 9H), 1.80-1.23 (m, 6H), 0.99 (s, 3H), 0.91 (s, 3H).

20

Example (Compound 459) NMR (CDCl₃): 7.39 (dd, 1H), 7.33-7.16 (m, 6H), 6.93 (d, 1H), 6.12 (s, 2H), 5.87 (m, 1H), 5.64 (d, 1H), 5.36 (d, 1H), 5.02 (q, 1H), 4.28 (br s, 1H), 4.08 (m, 1H), 4.00-3.62 (m, 5H), 3.35-2.70 (m, 9H), 2.00 (s, 3H), 1.70-1.25 (m, 8H), 0.96 (s, 3H), 0.91 (s, 3H).

30

Example (Compound 464) NMR (CDCl₃): 7.45 (s, 1H), 7.39 (dd, 1H), 7.31-7.15 (m, 6H), 7.11 (m, 1H), 6.93 (d, 1H),

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6.65-6.47 (m, 2H), 6.12 (s, 2H), 5.64 (d, 1H), 5.52 (d, 1H), 5.01 (q, 1H), 4.30 (br s, 1H), 4.09 (m, 1H), 4.00-3.57 (m, 5H), 3.45 (m, 2H), 3.30-2.97 (m, 4H), 2.95-2.70 (m, 3H), 1.80-1.25 (m, 8H), 0.97 (s, 3H), 0.90 (s, 3H).

5

Example (Compound 483) NMR (CDCl₃): 7.30 (dd, 1H), 7.25-7.08 (m, 6H), 6.85 (d, 1H), 6.59 (m, 1H), 6.03 (s, 2H), 5.59 (d, 1H), 5.30 (d, 1H), 4.94 (q, 1H), 4.21 (br s, 1H), 4.10-3.50 (m, 7H), 3.47-2.63 (m, 13H), 1.60-1.05 (m, 8H), 0.92 (s, 3H), 0.88 (s, 3H).

10

Example (Compound 494) NMR (CDCl₃): 7.28 (dd, 1H), 7.23-7.07 (m, 6H), 6.84 (d, 1H), 6.17-5.89 (m, 4H), 5.54 (d, 1H), 5.02 (d, 1H), 4.92 (q, 1H), 4.10-3.90 (m, 2H), 3.89-3.27 (m, 7H), 3.18-2.57 (m, 10H), 1.50 (m, 3H), 1.25 (m, 5H), 0.83 (d, 6H).

15

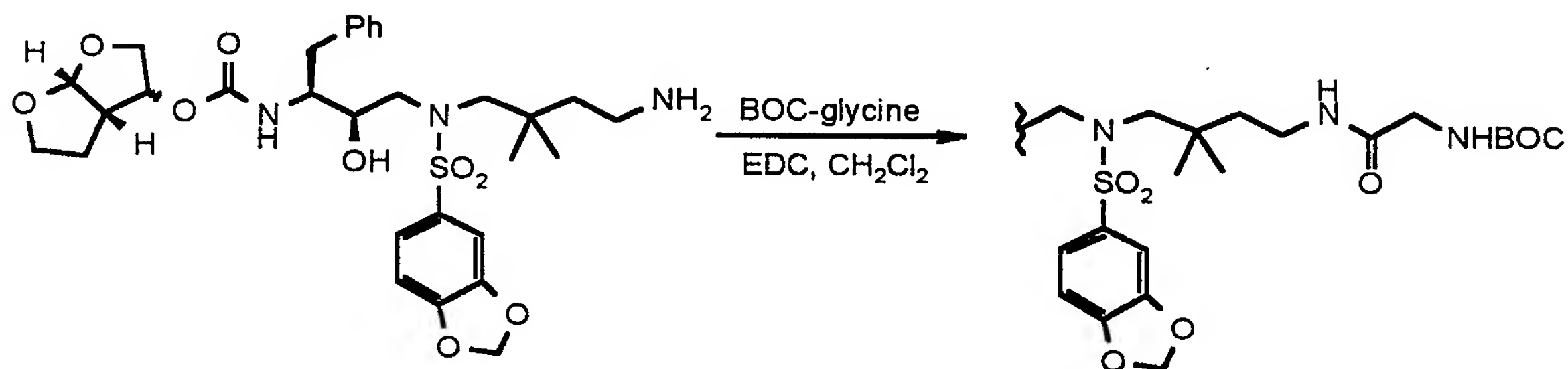
Example (Compound 496) NMR (CDCl₃): 7.28 (dd, 1H), 7.22-7.05 (m, 6H), 6.85 (d, 1H), 6.02 (s, 2H), 5.57 (d, 1H), 4.95 (m, 2H), 4.18 (br s, 1H), 3.94 (t, 1H), 3.89-3.22 (m, 10H), 3.18-2.60 (m, 7H), 1.70-1.08 (m, 14H), 0.86 (d, 6H).

20

Example (Compound 520)

25

Step 1:

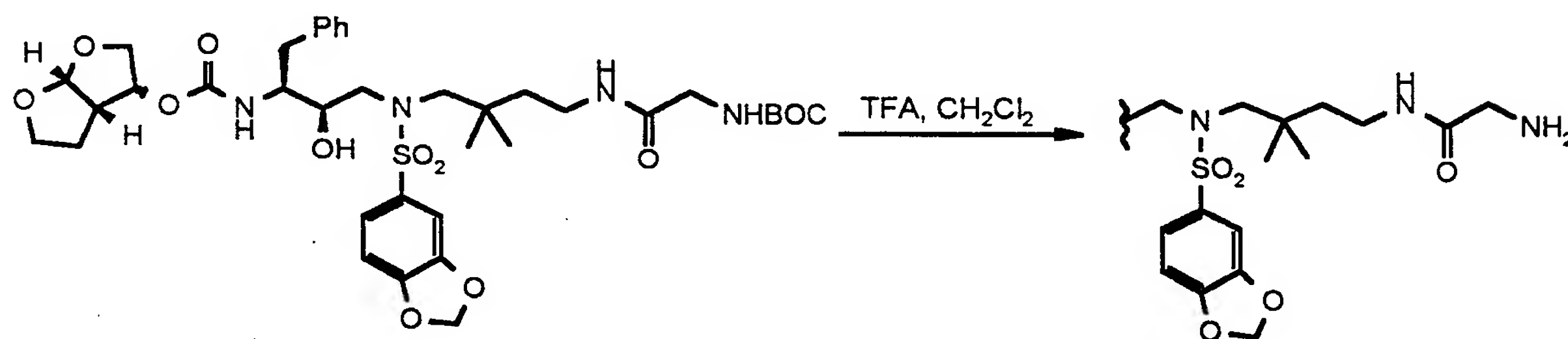


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(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(4-[(*tert*-butoxycarbonylamino)acetylamino]-2,2-
 dimethylbutyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-
 benzyl-2-hydroxypropylcarbamate

- 5 A solution of 0.300 g (0.484 mmol) of
 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(4-amino-2,2-dimethylbutyl)(1,3-benzodioxol-5-
 ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate
 (example 282), and 0.127 g (0.726 mmol) of BOC-glycine in
 10 8 mL of CH₂Cl₂ was treated with 0.139 g (0.726 mmol) of 1-
 ethyl-3-(dimethylaminopropyl)carbodiimide hydrochloride
 (EDC) and the resulting solution was stirred at RT.
 After 18 hours the solution was concentrated *in vacuo* and
 the crude material purified by flash chromatography (SiO₂,
 15 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.370 g (98%) of the
 desired compound as a white foam. MS (ESI): 799 (M+Na).

Step 2:



20

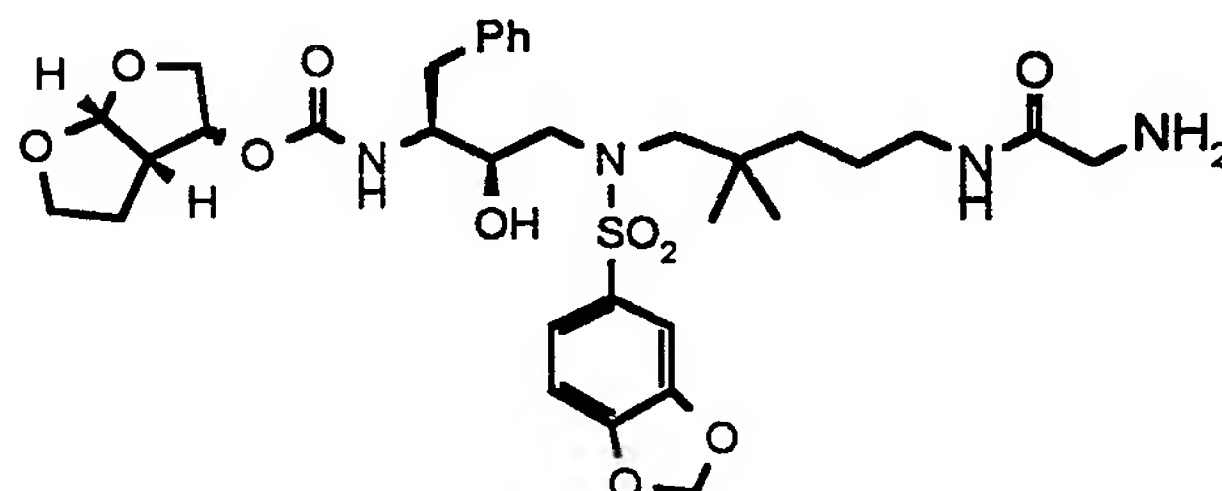
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(4-[(*tert*-butoxycarbonylamino)acetylamino]-2,2-
 dimethylbutyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-
 hydroxypropylcarbamate

- 25 A solution of 335 mg (0.431 mmol) of
 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(4-[(*tert*-butoxycarbonylamino)acetylamino]-2,2-
 dimethylbutyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-

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benzyl-2-hydroxypropylcarbamate in 30 mL of 1:1 TFA/CH₂Cl₂ was stirred at RT. After 2.5 hours the solution was concentrated to dryness and the residue redissolved in CH₂Cl₂. The solution was washed with 0.5N aqueous NaOH (1x), brine (2x), dried over MgSO₄, and concentrated *in vacuo* to afford 0.247 g (85%) of the desired compound as a white foam. NMR (CDCl₃): 7.62 (t, 1H), 7.38 (dd, 1H), 7.36-7.17 (m, 6H), 6.92 (d, 1H), 6.13 (s, 2H), 5.62 (d, 1H), 5.65 (d, 1H), 5.01 (q, 1H), 4.08-3.58 (m, 7H), 3.50-3.03 (m, 8H), 2.96-2.80 (m, 2H), 2.72 (dd, 1H), 2.30-1.66 (m, 3H), 1.62-1.38 (m, 2H), 1.26 (dd, 1H), 1.05 (s, 3H), 0.93 (s, 3H). MS(ESI): 677 (M+H).

Example (Compound 521)



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(5-[(amino)acetylamino]-2,2-dimethylpentyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate

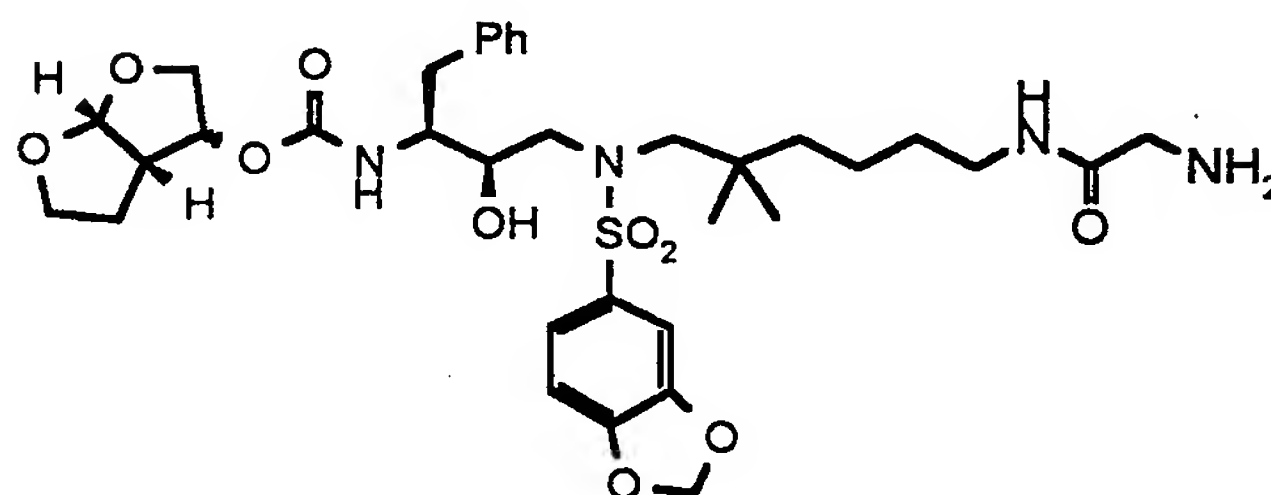
According to example 520, (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(5-amino-2,2-dimethylpentyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate (example 257) was converted to the title compound which was obtained as a white foam. NMR (CDCl₃): 7.49 (t, 1H), 7.40 (dd, 1H), 7.33-7.14 (m, 6H), 6.92 (d, 1H), 6.12 (s, 2H), 5.82 (d,

-321-

1H), 5.64 (d, 1H), 5.02 (q, 1H), 4.08-3.63 (m, 7H), 3.42 (s, 2H), 3.40-3.02 (m, 6H), 2.89 (m, 2H), 2.72 (dd, 1H), 2.61-2.00 (br, 2H), 1.53 (m, 3H), 1.34 (m, 3H), 1.00 (s, 3H), 0.92 (s, 3H). MS(ESI): 691(M+H).

5

Example (Compound 522)



10

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(6-[(amino)acetylamino]-2,2-dimethylhexyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate

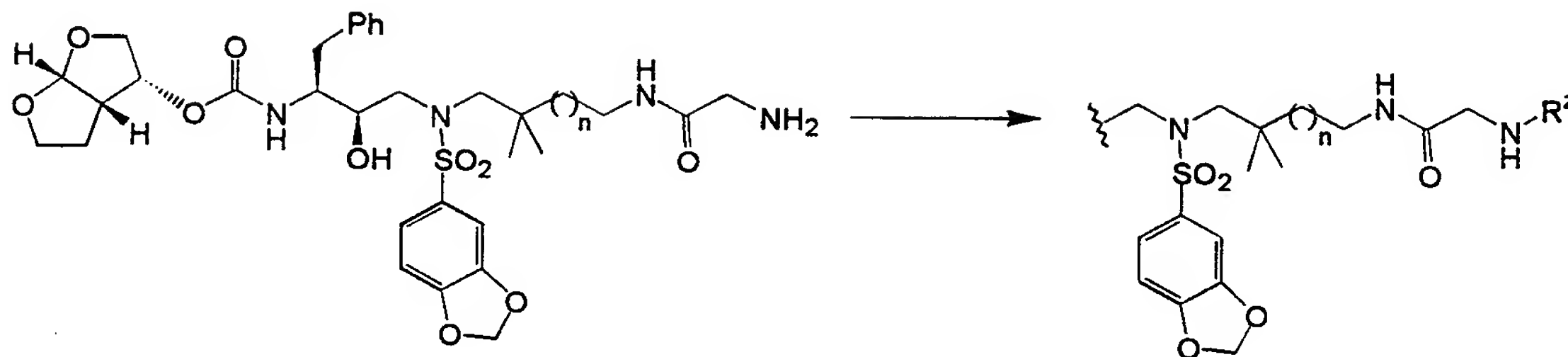
15 According to example 520, (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(6-amino-2,2-dimethylhexyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate (example 244) was converted to the title compound which was obtained as a

20 white foam. NMR (CDCl₃): 7.47-7.32 (m, 2H), 7.31-7.18 (m, 6H), 6.94 (d, 1H), 6.12 (s, 2H), 5.65 (d, 1H), 5.45 (d, 1H), 5.01 (q, 1H), 4.10 (t, 1H), 4.01-3.61 (m, 6H), 3.50-2.70 (m, 11H), 2.50-1.70 (br, 2H), 1.55 (m, 3H), 1.34 (m, 5H), 0.97 (s, 3H), 0.91 (s, 3H). MS(ESI): 705(M+H).

25

Examples (Compounds 523-543)

-322-



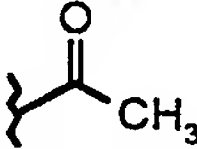
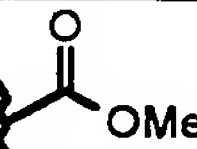
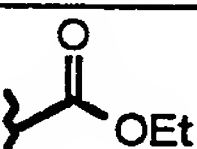
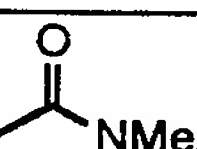
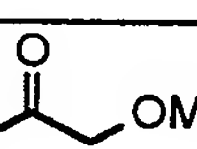

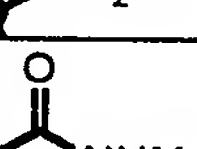
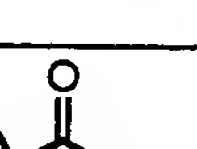

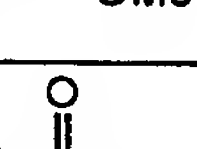
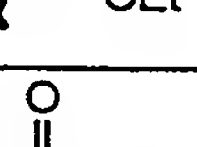
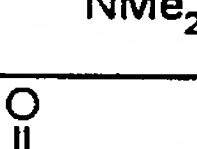

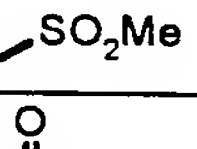
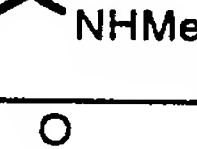
General procedure for reactions of amines 520, 521, and 522 with various electrophiles using scavenger resins:

- 5 The following reactions were carried out in parallel and were done in teflon-capped glass vials using a rotary shaker for agitation. To vials containing 130 mg of Amberlyst A-21 resin (Aldrich, prepared as in examples 373-519, not employed when methyl isocyanate was used as
- 10 the electrophile) were added 1 mL of 30 mM solutions of the amine starting materials in anhydrous THF. This was followed by addition of 0.15 mL (5 equiv.) of 1M solutions of electrophiles (acid chlorides, alkyl chloroformates, *N,N*-dimethylcarbonyl chloride,
- 15 methanesulfonyl chloride, methyl isocyanate) in anhydrous CH_2Cl_2 . After the resulting mixtures were shaken at RT for 18 hours, they were diluted with CH_2Cl_2 (3x), treated with 125 mg of PS-Trisamine resin (3.75 mmol/g, Argonaut Technologies, prepared as in examples 373-519, not
- 20 employed when methyl isocyanate was used as the electrophile) and shaken for an additional 18 hours. The resins were removed by filtration and the filtrates evaporated under a stream on nitrogen to afford the desired compounds.

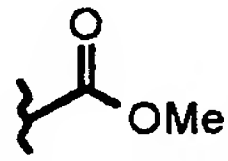
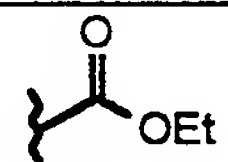
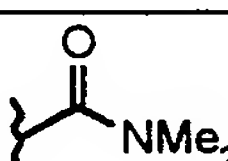
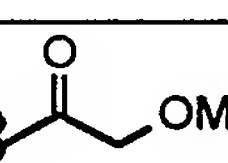
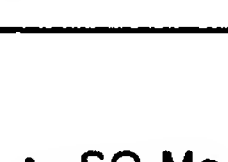
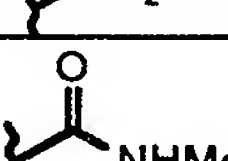
25

Mass Spectral Data for Compounds 523-543:

-323-

Example	Amine Precursor	N	R ^z	MS (ESI)
523	520	1		719 (M+H)
524	520	1		757 (M+Na)
525	520	1		749 (M+H)
526	520	1		770 (M+Na)
527	520	1		749 (M+H)
528	520	1		777 (M+Na)
529	520	1		756 (M+Na)
530	521	2		733 (M+H)
531	521	2		771 (M+Na)
532	521	2		785 (M+Na)
533	521	2		784 (M+Na)
534	521	2		763 (M+H)
535	521	2		791 (M+Na)
536	521	2		770 (M+Na)
537	522	3		747 (M+H)

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538	522	3		785 (M+Na)
539	522	3		777 (M+H)
540	522	3		798 (M+Na)
541	522	3		777 (M+H)
542	522	3		805 (M+Na)
543	522	3		784 (M+Na)

Proton NMR data for selected compounds from the above table:

5 Example (Compound 523) NMR (CDCl₃): 7.29 (dd, 1H), 7.23-7.04 (m, 6H), 6.81 (m, 3H), 6.02 (s, 2H), 5.72 (d, 1H), 5.55 (d, 1H), 4.96 (m, 1H), 4.03-3.47 (m, 8H), 3.29-2.54 (m, 10H), 2.10-1.40 (m, 5H), 1.32 (m, 1H), 1.10 (dd, 1H), 0.96 (s, 3H), 0.82 (s, 3H).

10

Example (Compound 524) NMR (CDCl₃): 7.29 (dd, 1H), 7.23-7.04 (m, 6H), 6.85 (d, 1H), 6.53 (m, 1H), 6.03 (s, 2H), 5.76 (m, 1H), 5.53 (m, 2H), 4.93 (q, 1H), 4.23 (br s, 1H), 3.98 (t, 1H), 3.87-3.44 (m, 10H), 3.38-3.02 (m, 4H), 15 3.01-2.71 (m, 3H), 2.62 (m, 2H), 1.80-1.25 (m, 3H), 1.16 (m, 1H), 0.99 (s, 3H), 0.84 (s, 3H).

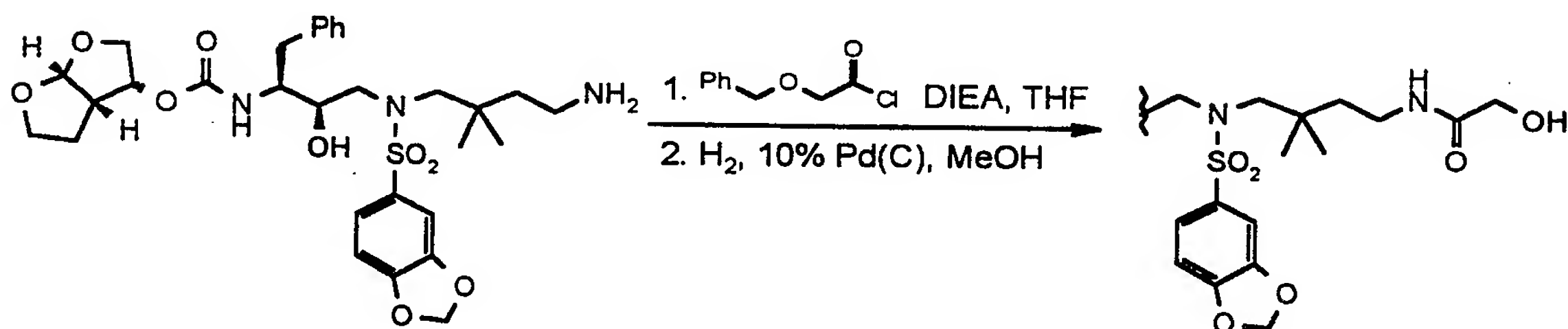
Example (Compound 531) NMR (CDCl₃): 7.28 (dd, 1H), 7.23-7.04 (m, 6H), 6.84 (d, 1H), 6.45 (m, 1H), 6.03 (s, 2H), 20 5.60-5.40 (m, 3H), 4.92 (q, 1H), 4.11 (d, 1H), 3.98 (m,

-325-

1H), 3.89-3.51 (m, 10H), 3.29-2.88 (m, 6H), 2.79 (m, 2H), 2.62 (dd, 1H), 1.60-1.19 (m, 6H), 0.91 (s, 3H), 0.82 (s, 3H).

5 Example (Compound 542) NMR (CDCl₃): 7.41-7.20 (m, 7H), 6.96 (d, 1H), 6.62 (m, 1H), 6.14 (s, 2H), 5.73 (t, 1H), 5.67 (d, 1H), 5.31 (d, 1H), 5.05 (q, 1H), 4.43 (d, 1H), 4.11 (m, 1H), 3.98-3.61 (m, 6H), 3.52 (m, 1H), 3.34-2.85 (m, 11H), 2.76 (dd, 1H), 1.70-1.25 (m, 8H), 1.01 (s, 3H),
10 0.90 (s, 3H).

Example (Compound 544)



15

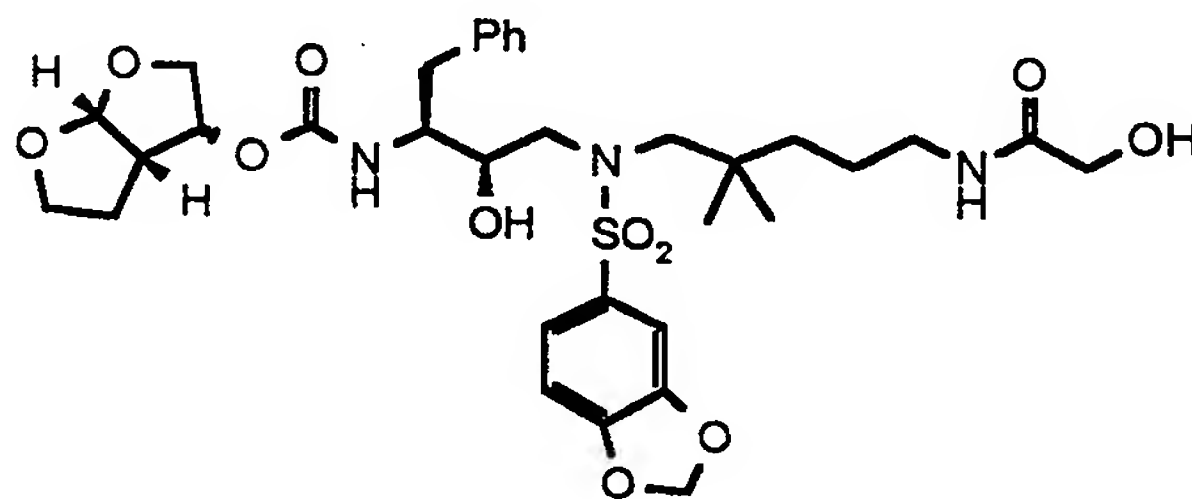
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
[(4-[(hydroxy)acetylamino]-2,2-dimethylbutyl)(1,3-
20 benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-
hydroxypropylcarbamate

A solution of 50 mg (0.080 mmol) of
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
[(4-amino-2,2-dimethylbutyl)(1,3-benzodioxol-5-
25 ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate
(example 282) and 21 μ L (0.12 mmol) of N,N-
diisopropylethylamine in 4 mL of anhydrous THF at 0°C was
treated with 13.3 μ L (0.084) of benzyloxyacetyl chloride.

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The resulting solution was allowed to warm to RT with stirring. After 2 hours the solution was concentrated in vacuo and the residue purified by flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH) to afford 52 mg (85%) of the
 5 desired intermediate as a white foam (m/e=768, M+H). This material (44 mg, 0.57 mmol) was then dissolved in 5 mL of MeOH and subjected to balloon hydrogenation in the presence of 25 mg of 10% Pd(C). After 2 hours the reaction vessel was purged with nitrogen, catalyst
 10 removed by filtration through celite, and the filtrate concentrated in vacuo to afford 31 mg (79%) of the desired product as a white foam. NMR (CDCl₃): 7.31 (dd, 1H), 7.26-7.07 (m, 6H), 6.92 (br s, 1H), 6.87 (d, 1H), 6.04 (s, 2H), 5.69 (d, 1H), 5.56 (d, 1H), 4.95 (m, 1H),
 15 4.12-3.90 (m, 3H), 3.86-3.62 (m, 4H), 3.54 (m, 1H), 3.36-3.15 (m, 5H), 3.02 (dd, 1H), 2.93 (d, 1H), 2.77 (q, 1H), 2.70-2.57 (m, 3H), 1.72 (m, 1H), 1.49 (m, 1H), 1.30 (m, 1H), 1.16-0.97 (m, 4H), 0.91 (s, 3H). MS(ESI): 678 (M+H).

20

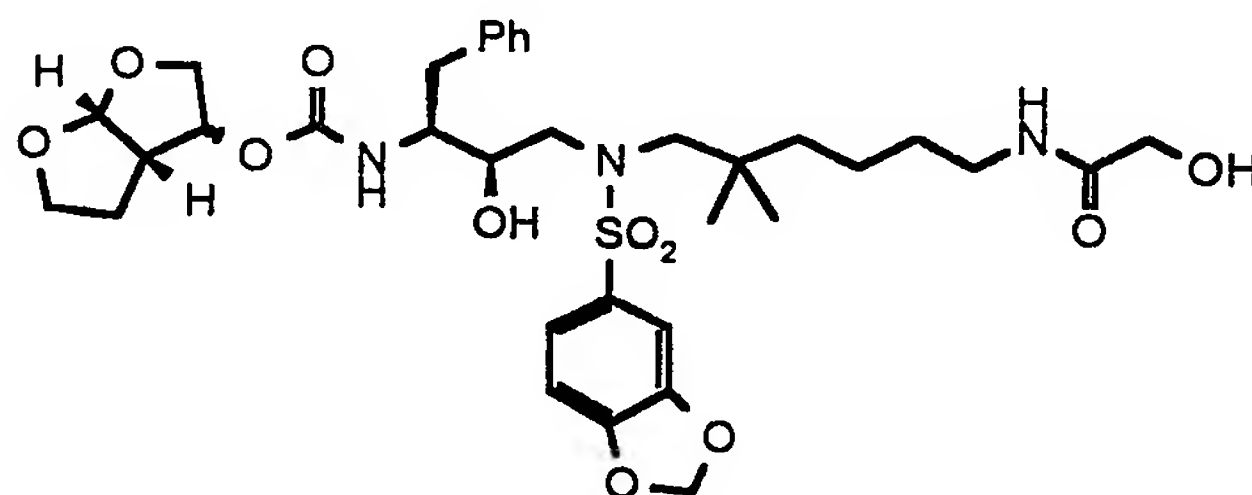
Example (Compound 545)

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-3-
 25 [(5-[(hydroxy)acetyl amino]-2,2-dimethylpentyl)(1,3-benzodioxol-5-yl)sulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate

-327-

According to example 544 , (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(5-amino-2,2-dimethylpentyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate was converted to the
 5 title compound which was obtained as a white foam. NMR (CDCl₃): 7.30 (dd, 1H), 7.27-7.10 (m, 6H), 6.88 (m, 2H), 6.05 (s, 2H), 5.55 (m, 2H), 4.96 (q, 1H), 4.14-3.91 (m, 3H), 3.89-3.71 (m, 3H), 3.70-3.54 (m, 2H), 3.36-2.90 (m, 8H), 2.81 (m, 2H), 2.63 (t, 1H), 1.50 (m, 3H), 1.41-1.17
 10 (m, 3H), 0.91 (s, 3H), 0.85 (s, 3H). MS(ESI): 692(M+H).

Example (Compound 546)



15

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(6-[(hydroxy)acetylamino]-2,2-dimethylhexyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate

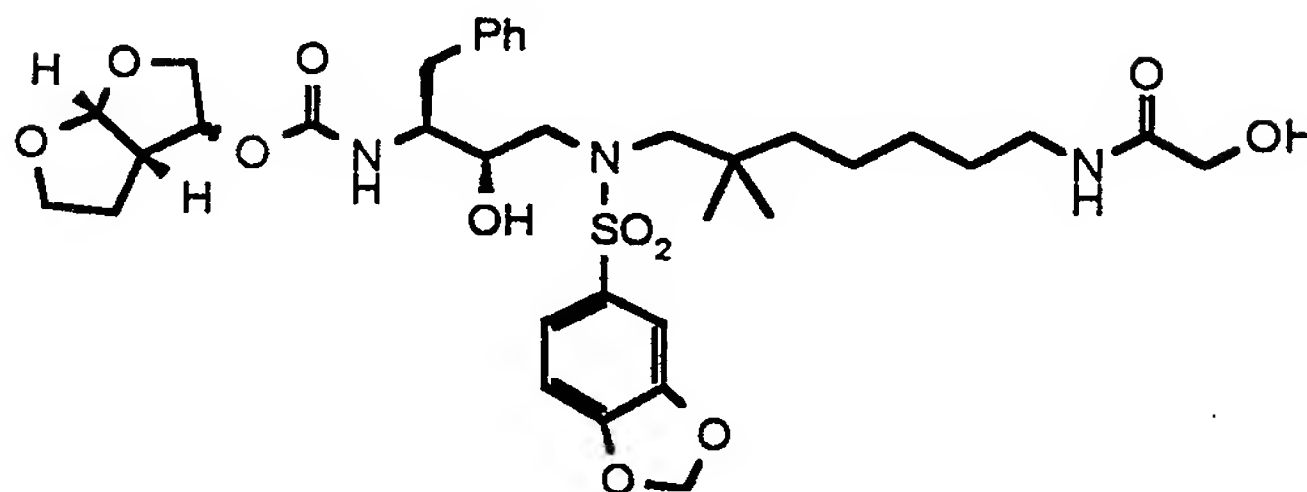
20 According to example 544, (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(6-amino-2,2-dimethylhexyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate was converted to the title compound which was obtained as a white foam. NMR
 25 (CDCl₃): 7.30 (dd, 1H), 7.27-7.11 (m, 6H), 6.88 (d, 1H), 6.74 (br s, 1H), 6.07 (s, 2H), 5.58 (d, 1H), 5.29 (d, 1H), 4.95 (q, 1H), 4.15-3.96 (m, 3H), 3.90-3.71 (m, 3H), 3.70-3.52 (m, 2H), 3.38 (m, 1H), 3.21 (m, 1H), 3.16-2.60

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(m, 9H), 1.51 (m, 3H), 1.29 (m, 5H), 0.91 (s, 3H), 0.83 (s, 3H). MS(ESI): 706(M+H).

Example (Compound 547)

5



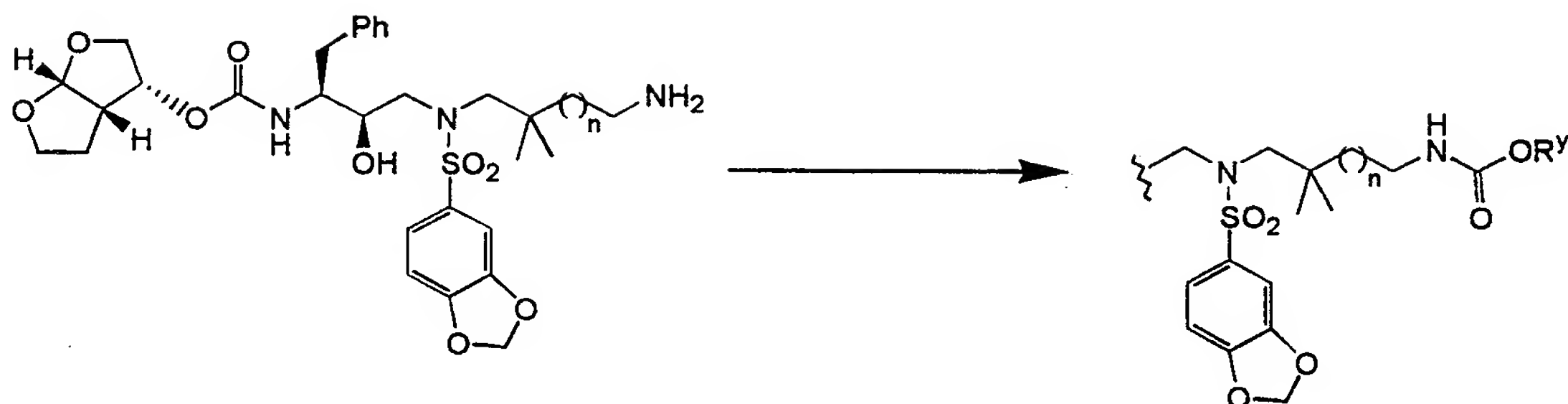
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(7-[(hydroxy)acetylamino]-2,2-dimethylheptyl)(1,3-
 10 benzodioxol-5-ylsulfonyl)amino]-1-benzyl-2-
 hydroxypropylcarbamate

According to example 544, (3R,3aS,6aR)hexahydrofuro[2,3-
 b]furan-3-yl N-(1S,2R)-3-[(7-amino-2,2-
 dimethylheptyl)(1,3-benzodioxol-5-ylsulfonyl)amino]-1-
 15 benzyl-2-hydroxypropylcarbamate was converted to the
 title compound which was obtained as a white foam. NMR
 (CDCl₃): 7.30 (dd, 1H), 7.26-7.08 (m, 6H), 6.88 (d, 1H),
 6.74 (br s, 1H), 6.06 (s, 2H), 5.58 (d, 1H), 5.20 (d,
 1H), 4.92 (q, 1H), 4.02 (br s, 3H), 3.92-3.71 (m, 3H),
 20 3.70-3.55 (m, 2H), 3.37-2.60 (m, 11H), 1.51 (m, 3H),
 1.40-1.11 (m, 7H), 0.89 (s, 3H), 0.85 (s, 3H). MS(ESI):
 720(M+H).

25

Examples (Compounds 548-555)

-329-

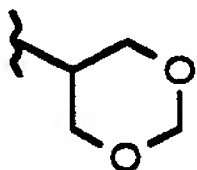


According to example 245, step 5 (with the exception that N-succinimidyl (3S)-tetrahydro-3-furanyl carbonate or 1,3-dioxan-5-yl 4-nitrophenyl carbonate was used as the acylating reagent), primary amines 282, 257, 244, and 283 were acylated to afford the desired products. See US 5,585,397 for synthesis of the above active carbonates.

10 Mass Spectral Data for Compounds 548-555:

Example	Amine Precursor	n	R ^y	MS (ESI)
548	282	1		756 (M+Na)
549	282	1		750 (M+H)
550	257	2		748 (M+H)
551	257	2		786 (M+Na)
552	244	3		784 (M+Na)
553	244	3		778 (M+H)
554	283	4		776 (M+H)

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555	283	4		792 (M+H)
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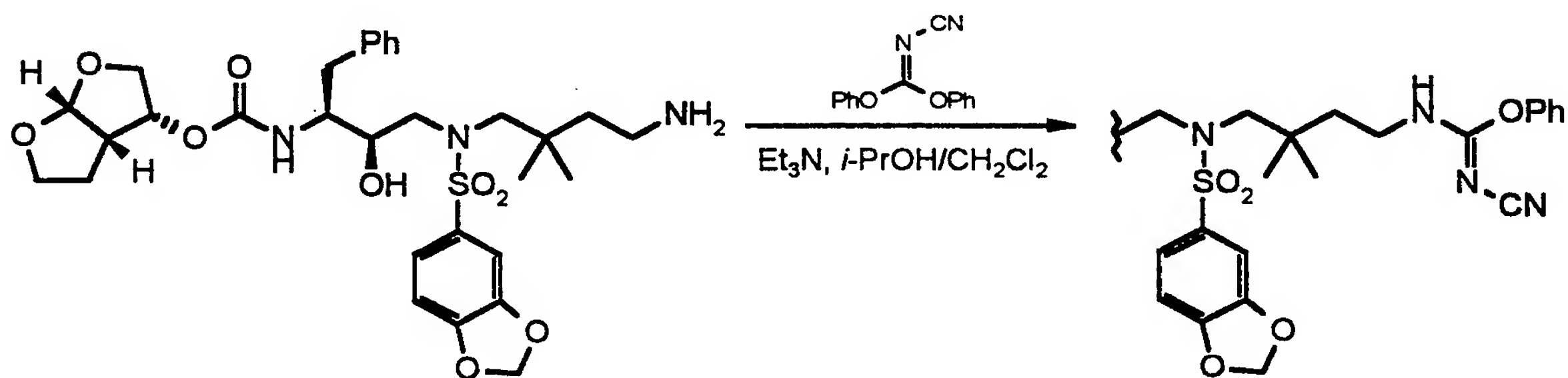
Proton NMR data for selected compounds from the above table:

5 Example (Compound- 548) NMR (CDCl₃): 7.37 (dd, 1H), 7.35-7.16 (m, 6H), 6.95 (d, 1H), 6.14 (s, 2H), 5.67 (d, 1H), 5.32-5.10 (m, 3H), 5.03 (q, 1H), 4.20-3.60 (m, 11H), 3.34-2.70 (m, 9H), 2.28-1.96 (m, 2H), 1.80-1.35 (m, 4H), 1.06 (s, 3H), 1.00 (s, 3H).

10

Example (Compound 553) NMR (CDCl₃): 7.31 (dd, 1H), 7.28-7.10 (m, 6H), 6.87 (d, 1H), 6.03 (s, 2H), 5.58 (d, 1H), 5.08 (m, 2H), 5.00-4.86 (m, 2H), 4.73 (d, 1H), 4.59 (m, 1H), 4.10-3.70 (m, 8H), 3.61 (m, 2H), 3.20-2.62 (m, 10H),
 15 1.60-1.17 (m, 8H), 0.84 (d, 6H).

Example (Compound 556)



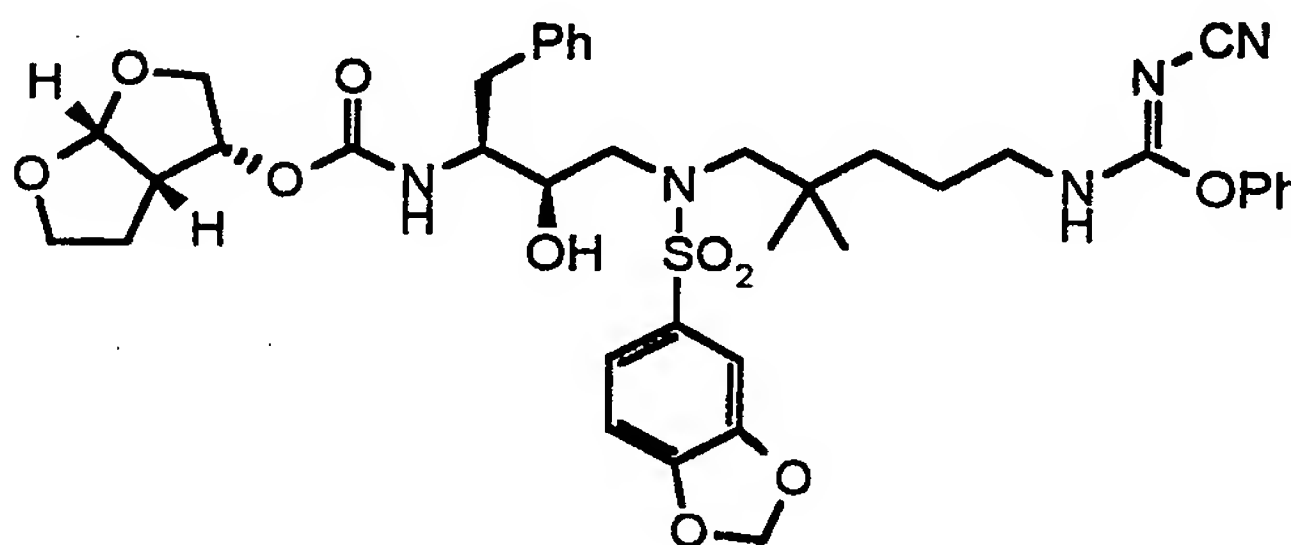
20

-331-

(3R, 3aS, 6aR)hexahydrofuro[2,3-b]furan-3-yl (1S, 2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl) (4-
 {[(cyanoimino) (phenoxy)methyl]amino}-2,2-
 dimethylbutyl)amino]-1-benzyl-2-hydroxypropylcarbamate

5 A solution of 0.250 g (0.403 mmol) of
 (3R, 3aS, 6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S, 2R)-3-
 [(4-amino-2,2-dimethylbutyl) (1,3-benzodioxol-5-
 ylsulfonyl)amino]-1-benzyl-2-hydroxypropylcarbamate in 15
 mL of 3:1 i-PrOH/CH₂Cl₂ was treated with 84 μ L (0.605
 10 mmol) of Et₃N followed by 0.106 g (0.443 mmol) of
 diphenyl cyanocarbonimide and the resulting solution
 was stirred at RT. After 3 hours the solution was
 concentrated in vacuo and the residue subjected to flash
 chromatography (SiO₂, 97:3 CH₂Cl₂/MeOH) to afford 0.269 g
 .15 (87%) of the desired compound as a white foam. NMR
 (CDCl₃): 7.40-7.10 (m, 9H), 7.06 (d, 2H), 6.91-6.69 (m,
 2H), 6.04 (s, 2H), 5.81-4.73 (complex pattern, rotamers,
 3H total), 4.06 (br s, 1H), 3.92-3.21 (m, 7H), 3.20-2.79
 (m, 6H), 2.69 (dd, 1H), 2.40-1.43 (m, 5H), 1.37 (m, 1H),
 20 0.95 (m, 6H). MS (ESI): 764 (M+H).

Example (Compound 557)



25

(3R, 3aS, 6aR)hexahydrofuro[2,3-b]furan-3-yl (1S, 2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl) (5-

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{ [(cyanoimino) (phenoxy)methyl] amino} -2,2-

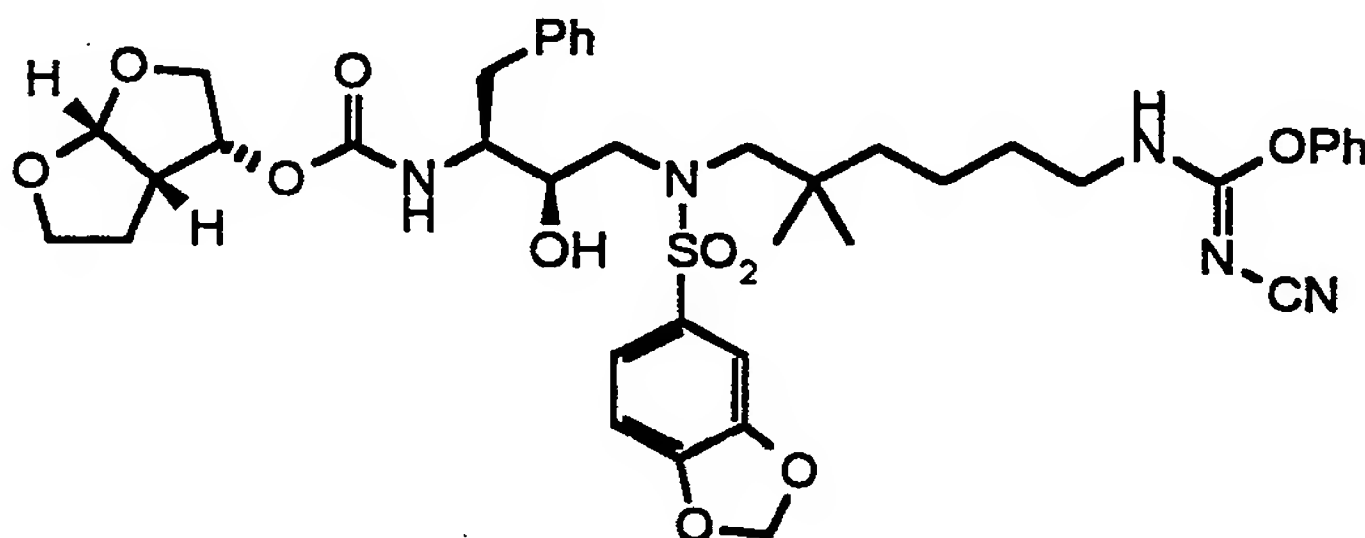
dimethylpentyl) amino] -1-benzyl-2-hydroxypropylcarbamate

According to example 556 , (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(5-amino-2,2-

5 dimethylpentyl) (1,3-benzodioxol-5-ylsulfonyl) amino] -1-benzyl-2-hydroxypropylcarbamate was converted to the title compound which was obtained as a white foam. NMR (CDCl₃): 7.40-7.02 (m, 11H), 6.88 (d, 1H), 6.54 (br s, 1H), 6.07 (s, 2H), 5.80-4.80 (complex pattern, rotamers, 3H total), 4.13-3.92 (m, 1H), 3.90-3.72 (m, 3H), 3.70-3.54 (m, 2H), 3.50-3.25 (m, 2H), 3.17-2.76 (m, 6H), 2.70 (dd, 1H), 2.40-1.18 (m, 8H), 0.90 (m, 6H). MS (ESI): 778 (M+H).

15

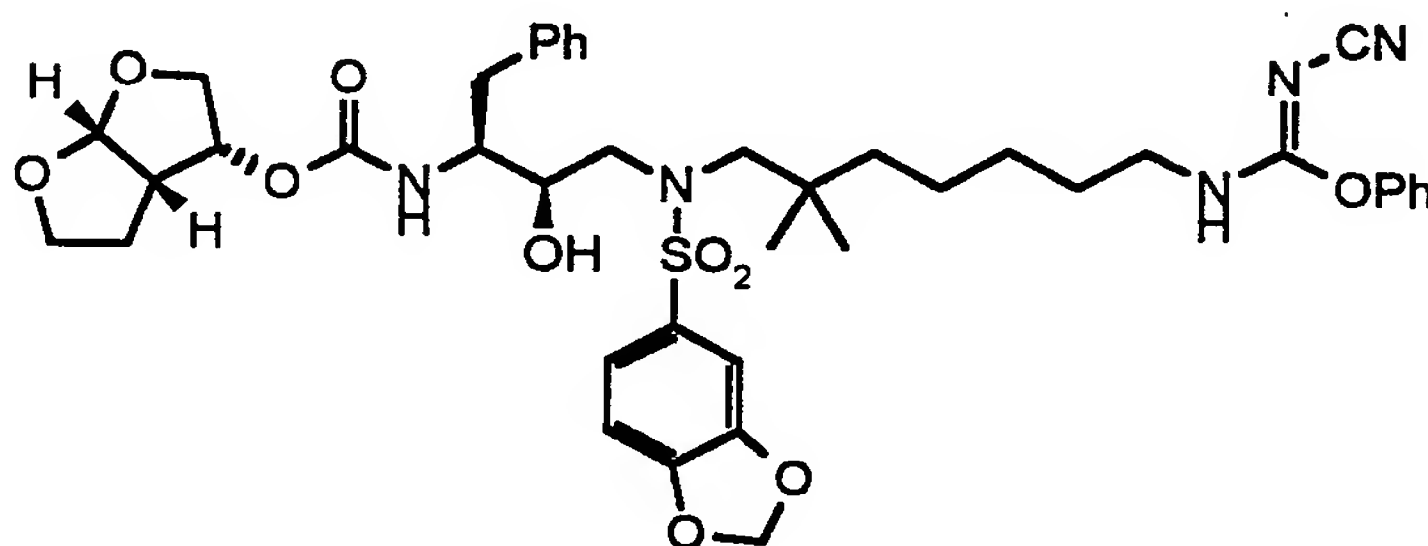
Example (Compound 558)



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 20 [(1,3-benzodioxol-5-ylsulfonyl) (6-
 { [(cyanoimino) (phenoxy)methyl] amino} -2,2-
 dimethylhexyl) amino] -1-benzyl-2-hydroxypropylcarbamate
 According to example 556, (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-[(6-amino-2,2-
 25 dimethylhexyl) (1,3-benzodioxol-5-ylsulfonyl) amino] -1-benzyl-2-hydroxypropylcarbamate was converted to the title compound which was obtained as a white foam. NMR

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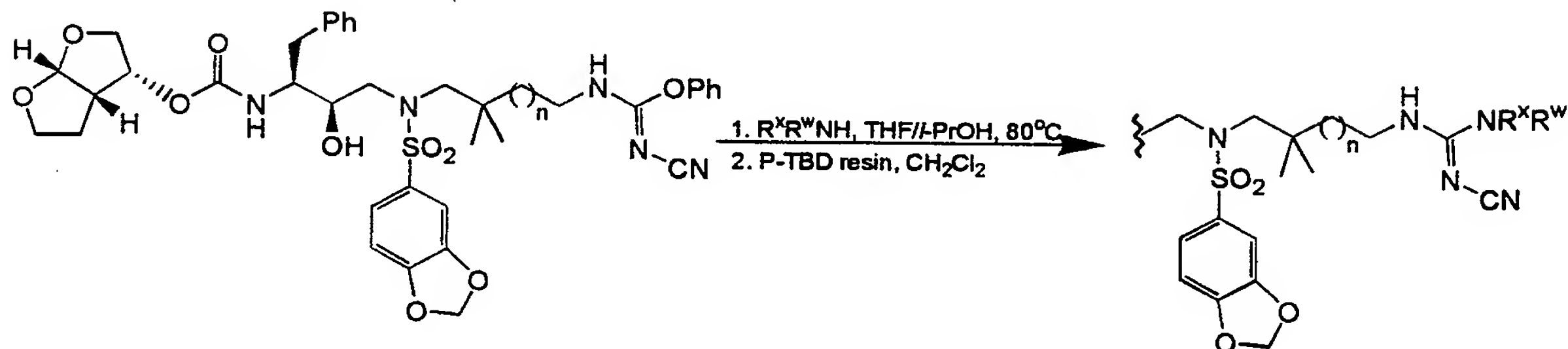
(CDCl₃): 7.40-7.01 (m, 11H), 6.87 (d, 1H), 6.52 (br s, 1H), 6.05 (s, 2H), 5.65-4.85 (complex pattern, rotamers, 3H total), 4.00 (m, 1H), 3.94-3.72 (m, 3H), 3.64 (m, 2H), 3.50-3.28 (m, 2H), 3.20-2.63 (m, 7H), 2.60-1.80 (br, 2H),
 5 1.70-1.45 (m, 3H), 1.43-1.20 (m, 5H), 0.89 (m, 6H).
 MS (ESI): 792 (M+H).

Example (Compound 559)

10

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl) (7-
 [(cyanoimino) (phenoxy)methyl]amino)-2,2-
 15 dimethylheptyl)amino]-1-benzyl-2-hydroxypropylcarbamate
 According to example 556, (3R,3aS,6aR)hexahydrofuro[2,3-
 b]furan-3-yl N-(1S,2R)-3-[(7-amino-2,2-
 dimethylheptyl) (1,3-benzodioxol-5-ylsulfonyl)amino]-1-
 benzyl-2-hydroxypropylcarbamate was converted to the
 20 title compound which was obtained as a white foam. NMR
 (CDCl₃): 7.45-7.00 (m, 11H), 6.88 (d, 1H), 6.34 (br s, 1H), 6.06 (s, 2H), 5.65-4.70 (complex pattern, rotamers, 3H total), 4.01 (m, 1H), 3.94-3.72 (m, 3H), 3.62 (m, 2H), 3.48-2.25 (m, 2H), 3.20-2.62 (7H), 2.60-1.75 (br, 2H),
 25 1.71-1.46 (m, 3H), 1.45-1.14 (m, 7H), 0.85 (m, 6H).
 MS (ESI): 806 (M+H).

-334-

Examples (Compounds 560-575)

- 5 Procedure for reactions of compounds 556-559 with various amines using P-TBD resin to remove phenol by-product:
The following reactions were carried out in parallel and were done in teflon-capped sealed tubes equipped with magnetic stirrers. Solutions of 0.026 mmol of starting materials 556, 557, 558, and 559 in 3 mL of 1:1 THF/i-PrOH were treated with excess (40 equiv.) amine. Ammonia and MeNH₂ were added as 2 M solutions in MeOH and EtOH respectively; Me₂NH was added as a 2M solution in THF; and n-PrNH₂ was added neat. The vessels were capped and solutions heated to 80°C with stirring. After 3 hours the vessels were cooled to RT and examined by tlc which indicated all reactions to be complete. The solutions were cooled to RT and evaporated under a stream of nitrogen. The resulting residues were dissolved in 3 mL of CH₂Cl₂ and the solutions treated with 100 mg of polymer bound 1,3,4,6,7,8-hexahydro-2H-pyrimido [1,2-a]pyrimidine (P-TBD resin, 2.6 mmol/g, Aldrich). The mixtures were shaken on a rotary shaker for 3 hours, filtered to remove the resin, and the filtrates evaporated under a stream of nitrogen to afford the desired products.

Mass Spectral Data for Compounds 560-575:

-335-

Example	Starting Material	n	R ^x	R ^y	MS (ESI)
560	556	1	H	H	687 (M+H)
561	556	1	Me	H	701 (M+H)
562	556	1	n-Pr	H	729 (M+H)
563	556	1	Me	Me	715 (M+H)
564	557	2	H	H	701 (M+H)
565	557	2	Me	H	715 (M+H)
566	557	2	n-Pr	H	743 (M+H)
567	557	2	Me	Me	729 (M+H)
568	558	3	H	H	715 (M+H)
569	558	3	Me	H	729 (M+H)
570	558	3	n-Pr	H	757 (M+H)
571	558	3	Me	Me	743 (M+H)
572	559	4	H	H	729 (M+H)
573	559	4	Me	H	743 (M+H)
574	559	4	n-Pr	H	771 (M+H)

-336-

575	559	4	Me	Me	757 (M+H)
-----	-----	---	----	----	-----------

Proton NMR data for selected compounds from the above table:

5

Example (Compound 560) NMR (CDCl₃): 7.31 (dd, 1H), 7.24-7.02 (m, 6H), 6.88 (d, 1H), 6.02 (s, 2H), 5.70-5.40 (m, 3H), 4.92 (m, 1H), 4.05 (br s, 1H), 3.96-3.42 (m, 8H), 3.36-2.50 (m, 9H), 1.80-1.07 (m, 4H), 0.98 (s, 3H), 0.82 (s, 3H).

10

Example (Compound 565) NMR (CDCl₃): 7.40-7.04 (m, 7H), 6.88 (d, 1H), 6.07 (s, 2H), 5.58 (d, 1H), 5.45 (br s, 2H), 5.11 (d, 1H), 4.91 (m, 1H), 4.01 (m, 1H), 3.93-3.50 (m, 6H), 3.24-2.50 (m, 12H), 1.70-1.15 (m, 6H), 0.84 (d, 6H).

15

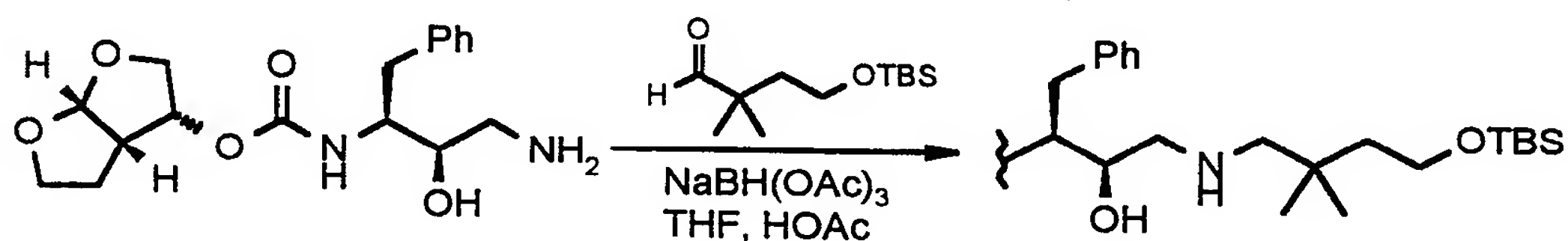
Example (Compound 571) NMR (CDCl₃): 7.32 (dd, 1H), 7.30-7.08 (m, 6H), 6.90 (d, 1H), 6.07 (s, 2H), 5.61 (d, 1H), 5.31 (d, 1H), 5.00 (m, 2H), 4.01 (t, 1H), 3.96-3.61 (m, 5H), 3.50 (m, 2H), 3.20-2.65 (m, 14H), 1.70-1.20 (m, 8H), 0.90 (d, 6H).

20

25

Example (Compound 576)

Step 1:



-337-

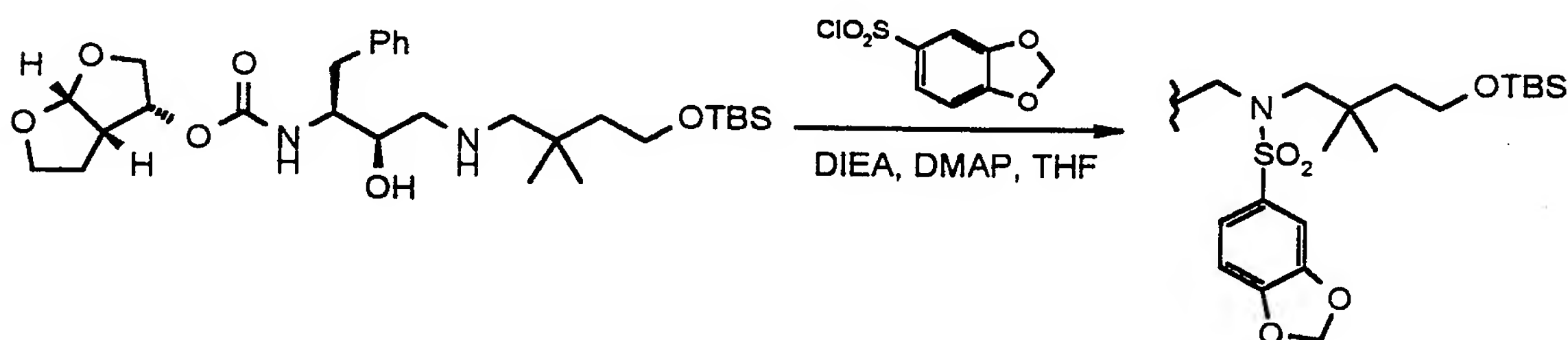
(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-1-benzyl-3-[(4-[*tert*-butyldimethylsiloxy]-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate

5 According to example 257, step 5,

(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-[(1*S*,2*R*)-3-amino-1-benzyl-2-hydroxypropyl]carbamate was subjected to reductive alkylation with of 4-(*tert*-butyldimethylsiloxy)-2,2-dimethylbutyraldehyde (example 221) to afford the desired compound as a white solid.

MS (ESI): 551 (M+H).

Step 2:



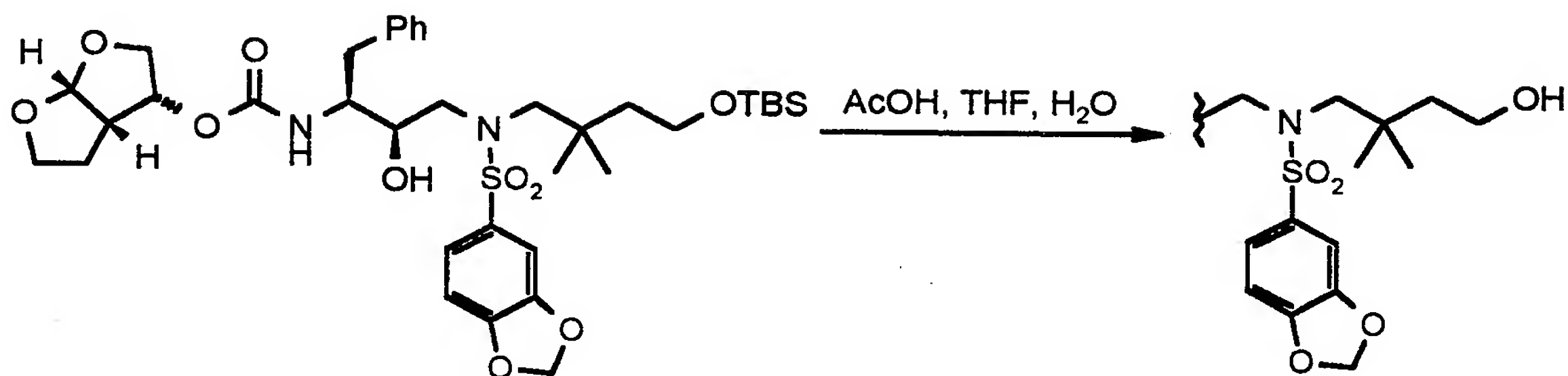
(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(4-[*tert*-butyldimethylsiloxy]-2,2-dimethylbutyl)amino]-1-benzyl-2-hydroxypropylcarbamate

According to example 257, step 6,

(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-1-benzyl-3-[(4-[*tert*-butyldimethylsiloxy]-2,2-dimethylbutyl)amino]-2-hydroxypropylcarbamate was subjected to sulfonylation to afford the desired sulfonamide as a white foam. MS (ESI): 757 (M+H).

Step 3:

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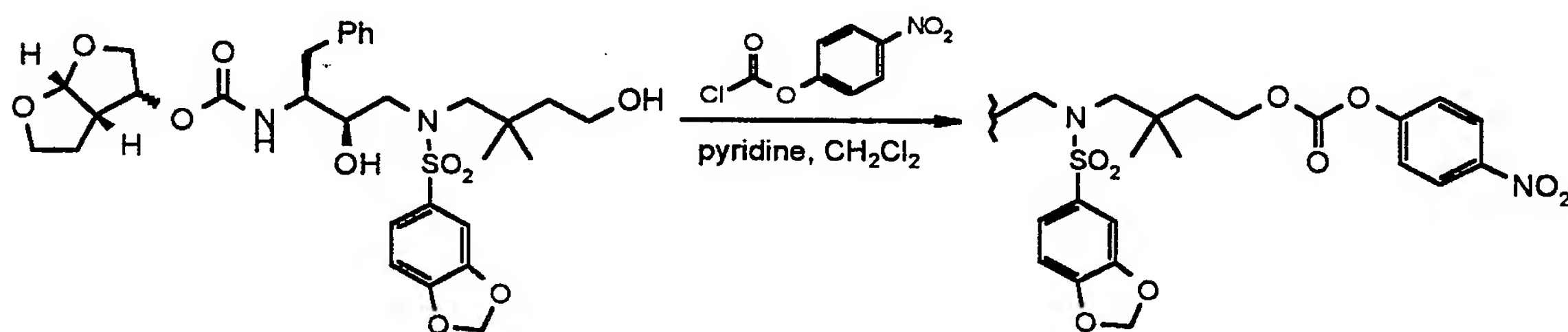


(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(2,2-dimethyl-4-
 5 hydroxybutyl)amino]-1-benzyl-2-hydroxypropylcarbamate

A solution of 0.67 g (0.91 mmol) of
 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(4-[tert-
 butyldimethylsiloxy]-2,2-dimethylbutyl)amino]-1-benzyl-2-
 10 hydroxypropylcarbamate in 25 mL of 3:1:1 AcOH/THF/H₂O was
 stirred at RT for 3 hours and then concentrated *in vacuo*.
 The crude product was purified by flash chromatography
 (SiO₂, hexane/EtOAc) to afford 0.50 g (89%) of the desired
 alcohol as a white foam. MS(ESI): 621(M+H).

15

Step 4:



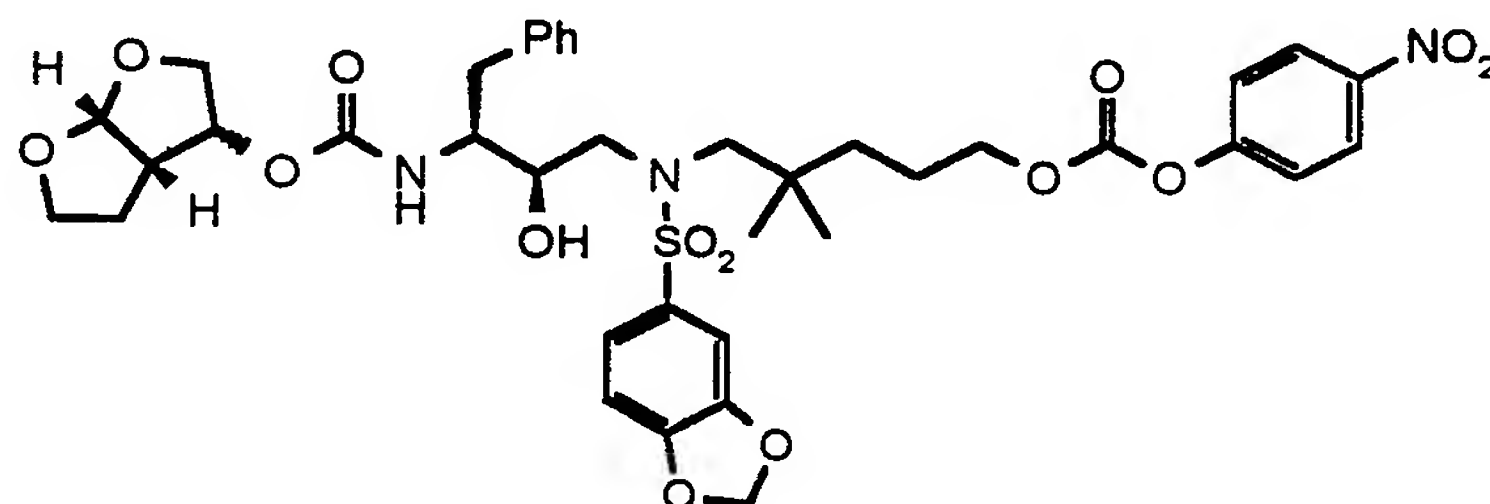
20 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(2,2-dimethyl-4-[(4-
 nitrophenyloxy)carbonyloxy]butyl)amino]-1-benzyl-2-
 hydroxypropylcarbamate

-339-

According to example 234, step1,
 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(2,2-dimethyl-4-
 hydroxybutyl)amino]-1-benzyl-2-hydroxypropylcarbamate was
 5 converted to the desired compound which was obtained as a
 white foam. NMR (CDCl₃): 8.22 (d, 2H), 7.32 (m, 3H), 7.21
 (m, 2H), 7.12 (m, 4H), 6.87 (d, 1H), 6.04 (s, 2H), 5.60
 (d, 1H), 4.95 (q, 1H), 4.84 (d, 1H), 4.35 (t, 2H), 4.02
 (m, 1H), 3.93-3.72 (m, 4H), 3.62 (m, 2H), 3.20-2.79 (m,
 10 6H), 2.69 (dd, 1H), 1.80 (m, 2H), 1.55 (m, 1H), 1.31 (m,
 1H), 1.01 (s, 6H). MS(ESI): 808(M+Na).

Example (Compound 577)

15



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl N-(1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(2,2-dimethyl-5-[(4-
 20 nitrophenoxy)carbonyloxy]pentyl)amino]-1-benzyl-2-
 hydroxypropylcarbamate

The title compound was prepared according to example 576
 with the exception that 5-(tert-butyldimethylsiloxy)-2,2-
 dimethylpentanal (see example 272) was used as the
 25 aldehyde component in step 1. NMR (CDCl₃): 8.23 (d, 2H),
 7.33 (m, 3H), 7.26-7.09 (m, 6H), 6.88 (d, 1H), 6.06 (s,
 2H), 5.59 (d, 1H), 4.95 (q, 1H), 4.85 (d, 1H), 4.26 (m,
 2H), 4.01 (m, 1H), 3.97-3.72 (m, 4H), 3.62 (m, 2H), 3.20-

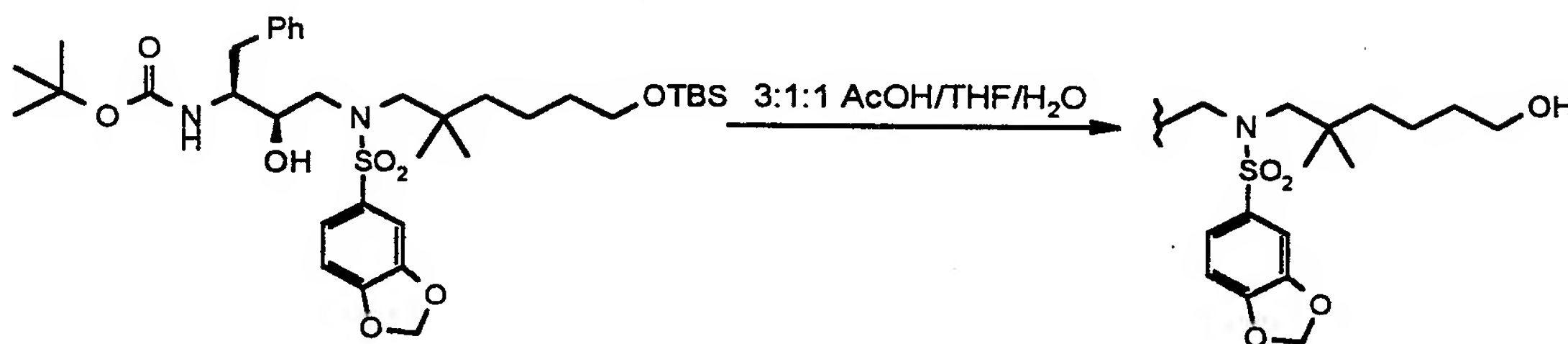
-340-

2.74 (m, 6H), 2.68 (dd, 1H), 1.71 (m, 2H), 1.52 (m, 1H),
1.38 (m, 3H), 0.92 (s, 6H). MS (ESI): 822 (M+Na).

Example (Compound 578)

5

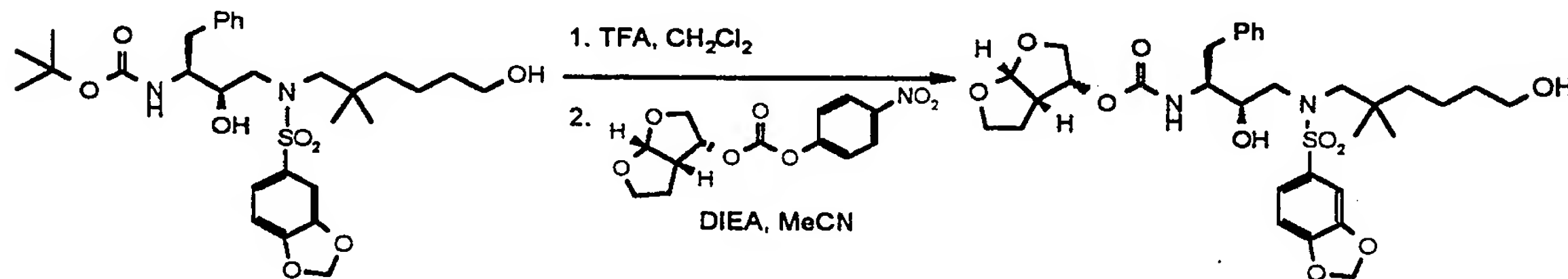
Step 1:



10 *tert*-Butyl *N*-(1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (2,2-dimethyl-6-hydroxyhexyl) amino]-1-benzyl-2-hydroxypropylcarbamate

According to example 576, step 3, *tert*-Butyl *N*-(1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (2,2-dimethyl-6-[*tert*-butyldimethylsiloxy]hexyl) amino]-1-benzyl-2-hydroxypropylcarbamate (prepared in a manner similar to
15 example 221) was desilylated to afford the desired compound as a white foam. MS (ESI): 593 (M+H).

Step 2:



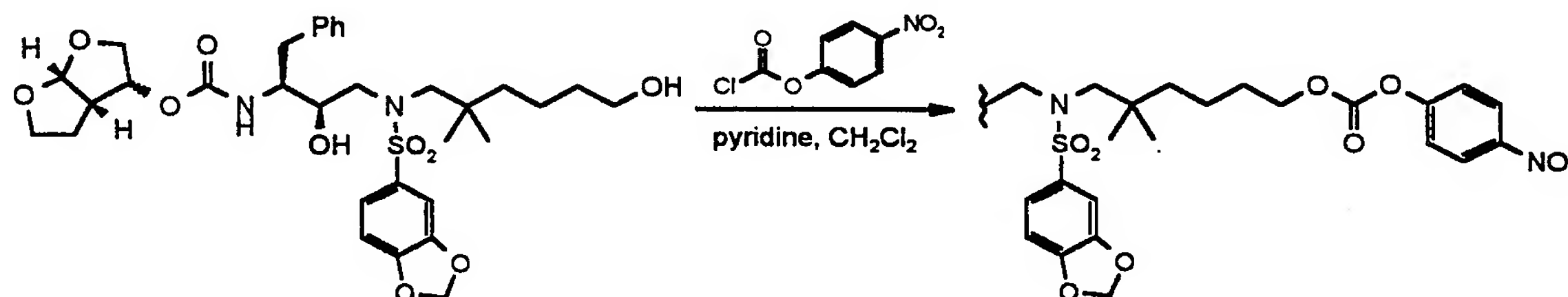
20

(3*R*,3*aS*,6*aR*) hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (2,2-dimethyl-6-hydroxyhexyl) amino]-1-benzyl-2-hydroxypropylcarbamate

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According to example 245, steps 4 and 5, *tert*-butyl *N*-(1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(2,2-dimethyl-6-hydroxyhexyl)amino]-1-benzyl-2-hydroxypropylcarbamate was converted to the desired product which was obtained as a white foam. MS(ESI): 671(M+Na).

Step 3:



10

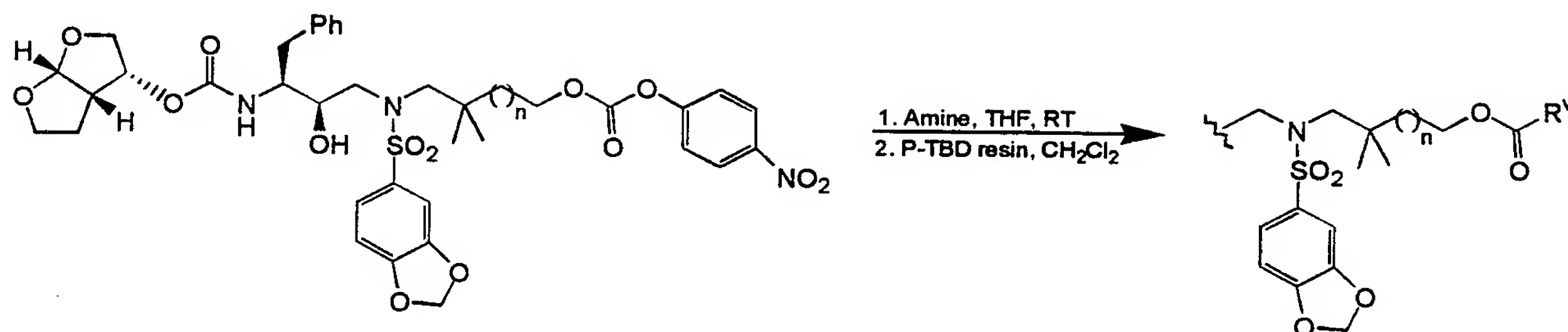
(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl *N*-(1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(2,2-dimethyl-6-[(4-nitrophenoxy)carbonyloxy]hexyl)amino]-1-benzyl-2-hydroxypropylcarbamate

15 The title compound was prepared according to example 234, step 1. NMR (CDCl₃): 8.23 (d, 2H), 7.32 (m, 3H), 7.27-7.08 (m, 6H), 6.86 (d, 1H), 6.06 (s, 2H), 5.58 (d, 1H), 4.96 (q, 1H), 4.83 (d, 1H), 4.25 (t, 2H), 4.01 (m, 1H), 3.91 (m, 2H), 3.80 (m, 2H), 3.61 (m, 2H), 3.20-2.73 (m, 6H), 2.70 (dd, 1H), 1.70 (m, 2H), 1.55 (m, 1H), 1.32 (m, 5H), 0.89 (d, 6H). MS(ESI): 836(M+H).

20

Examples (Compounds 579-604)

-342-



Procedure for reactions of compounds 576-578 with various amines using P-TBD resin to remove 4-nitrophenol by-product:

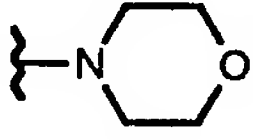


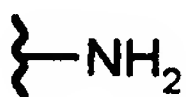
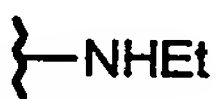




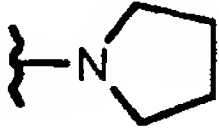
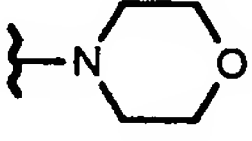

The following reactions were carried out in parallel and were done in teflon-capped vials using a rotary shaker for agitation. Solutions of 0.025 mmol of starting materials 576, 577, and 578 in 1 mL of anhydrous THF were treated with 10 equiv. of amine. Ammonia was added as a 2M solution in MeOH; ethylamine and dimethylamine were added as 2M solutions in THF; and the remaining amines were added neat. The solutions were shaken briefly and then allowed to stand at RT. After 2 hours tlc indicated all reactions to be complete. The solutions were evaporated under a stream on nitrogen. The residues were dissolved in 4 mL of CH₂Cl₂ and the solutions treated with 100 mg of polymer bound 1,3,4,6,7,8-hexahydro-2H-pyrimido [1,2-a]pyrimidine (P-TBD resin, 2.6 mmol/g, Aldrich). The mixtures were shaken for 3 hours, filtered to remove the resin, and the filtrates evaporated under a stream of nitrogen to afford the desired products.

Mass Spectral Data for Compounds 579-604:

-343-

Example	Starting Material	N	R ^v	MS (ESI)
579	576	1		692 (M+H)
580	576	1		664 (M+H)
581	576	1		692 (M+H)
582	576	1		706 (M+H)
583	576	1		706 (M+H)
584	576	1		720 (M+H)
585	576	1		704 (M+H)
586	576	1		718 (M+H)
587	576	1		734 (M+H)
588	576	1		679 (M+H)
589	577	2		742 (M+Na)
590	577	2		756 (M+Na)
591	577	2		740 (M+Na)
592	577	2		732 (M+H)

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593	577	2		748 (M+H)
594	577	2		693 (M+H)
595	578	3		720 (M+H)
596	578	3		692 (M+H)
597	578	3		720 (M+H)
598	578	3		734 (M+H)
599	578	3		734 (M+H)
600	578	3		748 (M+H)
601	578	3		732 (M+H)
602	578	3		746 (M+H)
603	578	3		762 (M+H)
604	578	3		707 (M+H)

Proton NMR data for selected compounds from the above table:

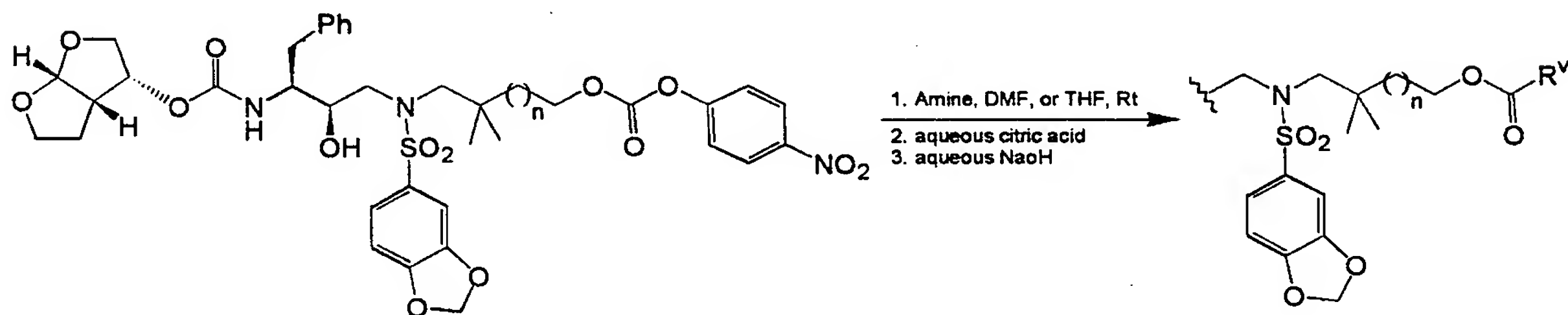
-345-

Example (Compound 579) NMR (CDCl₃): 7.30 (dd, 1H), 7.25-7.09 (m, 6H), 6.84 (d, 1H), 6.06 (s, 2H), 5.58 (d, 1H), 5.32 (d, 1H), 4.92 (q, 1H), 4.09 (m, 3H), 3.94 (t, 1H), 3.88 (dd, 1H), 3.84-3.53 (m, 5H), 3.24-2.97 (m, 4H), 2.83 (m, 7H), 2.66 (dd, 1H), 1.80-1.22 (m, 4H), 0.97 (s, 3H), 0.93 (s, 3H).

Example (Compound 589) NMR (CDCl₃): 7.39 (dd, 1H), 7.33-7.12 (m, 6H), 6.93 (d, 1H), 6.13 (s, 2H), 5.66 (d, 1H), 5.32 (d, 1H), 5.01 (m, 2H), 4.25-3.64 (m, 9H), 3.32-2.66 (m, 9H), 1.80-1.23 (m, 9H), 0.97 (m, 9H).

Example (Compound 596) NMR (CDCl₃): 7.32 (dd, 1H), 7.26-7.08 (m, 6H), 6.85 (d, 1H), 6.06 (s, 2H), 5.59 (d, 1H), 5.06 (d, 1H), 4.96 (q, 1H), 4.69 (br s, 2H), 4.17-3.55 (m, 9H), 3.20-2.64 (m, 7H), 1.75-1.19 (m, 8H), 0.88 (s, 6H).

20

Examples (Compounds 605-616)

Procedure for reactions of compounds 576-578 with various amines employing aqueous work-up:

The following reactions were carried out in parallel and were done in teflon-capped vials using a rotary shaker for agitation. Solutions of 0.038 mmol of starting

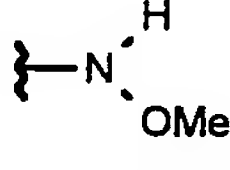
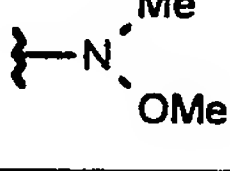
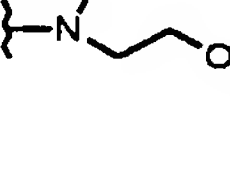
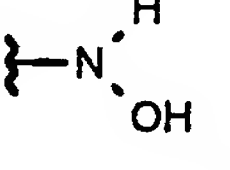
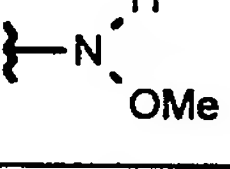
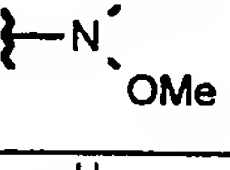
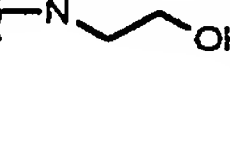
-346-

materials 576, 577, and 578 in 1 mL of anhydrous DMF (THF was used for examples 608, 612, and 616) were treated with 0.13 mL of *N,N*-diisopropylethylamine (omitted for examples 608, 612, and 616) followed by 15 equiv. of amine (hydroxylamine hydrochloride, methoxylamine hydrochloride, *N,O*-dimethylhydroxylamine hydrochloride, or ethanolamine). The resulting solutions were shaken at RT. After 18 hours the solutions were diluted with 5 mL of CH₂Cl₂, washed with 10% aqueous citric acid (3x), 0.5N aqueous NaOH (2x), aqueous brine (1x), and then dried over MgSO₄. The drying agent was removed by filtration and the filtrates evaporated under a stream of nitrogen. Products 605, 607, 609, 611, 613, and 615 were purified by preparative tlc (2 mm silica gel plate, 98:2 CH₂Cl₂/MeOH). The remaining products were found to be sufficiently pure for biological evaluation.

Mass Spectral Data for Compounds 605-616:

Example	Starting Material	N	R ^v	MS (ESI)
605	576	1		680 (M+H)
606	576	1		694 (M+H)
607	576	1		708 (M+H)
608	576	1		708 (M+H)
609	577	2		694 (M+H)

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610	577	2		708 (M+H)
611	577	2		722 (M+H)
612	577	2		722 (M+H)
613	578	3		708 (M+H)
614	578	3		722 (M+H)
615	578	3		736 (M+H)
616	578	3		736 (M+H)

Proton NMR data for selected compounds from the above table:

5

Example (Compound 606) NMR (CDCl₃): 8.01 (br s, 1H), 7.38 (dd, 1H), 7.33-7.17 (m, 6H), 6.93 (d, 1H), 6.14 (s, 2H), 5.64 (d, 1H), 5.18 (d, 1H), 5.02 (q, 1H), 4.36-4.12 (m, 2H), 4.05 (t, 1H), 3.98-3.62 (m, 8H), 3.30-2.61 (m, 8H), 1.85-1.22 (m, 4H), 1.04 (s, 3H), 1.00 (s, 3H).

10

Example (Compound 609) NMR (CDCl₃): 7.50-7.16 (m, 8H), 6.96 (d, 1H), 6.12 (s, 2H), 5.66 (d, 1H), 5.38 (d, 1H), 5.03 (m, 1H), 4.32-3.57 (m, 9H), 3.32-3.08 (m, 3H), 3.08-2.62 (m, 5H), 1.80-1.25 (m, 6H), 0.99 (s, 3H), 0.90 (s, 3H).

15

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Example (Compound 616) NMR (CDCl₃): 7.44-7.13 (m, 7H), 6.94 (d, 1H), 6.12 (s, 2H), 5.66 (d, 1H), 5.50-5.12 (m, 2H), 5.01 (q, 1H), 4.24-3.54 (m, 13H), 3.40-2.63 (m, 8H), 1.61 (m, 3H), 1.46-1.20 (m, 5H), 0.93 (s, 6H).

5

EXAMPLE

Anti-Viral Activity

10 We measured the enzyme inhibition constants of the compounds listed in Table I against HIV-1 protease using the methods of: B. Maschera et al., "Human Immunodeficiency Virus: Mutations in the Viral Protease that Confer Resistance to Saquinavir Increase the
15 Dissociation Rate Constant for the Protease-Saquinavir Complex", J. Biol. Chem., 271, pp. 33231-33235 (1996); and M. V. Toth et al., Int. J. Peptide Protein Res. 36, pp. 544-550. (1990)

Antiviral activity assay in MT4 cells

20 Antiviral HIV activity and compound-induced cytotoxicity were measured in parallel by means of a propidium iodide based procedure in the human T-cell lymphotropic virus transformed cell line MT4. Aliquots of the test compounds were serially diluted in medium
25 (RPMI 1640, 10% fetal calf serum (FCS), and gentamycin) in 96-well plates (Costar 3598) using a Cetus Pro/Pette. Exponentially growing MT4 cells were harvested and centrifuged at 1000 rpm for 10 min in a Jouan centrifuge (model CR 4 12). Cell pellets were resuspended in fresh
30 medium (RPMI 1640, 20% FCS, 20% IL-2, and gentamycin) to a density of 5×10^5 cells/ml. Cell aliquots were infected by the addition of HIV-1 (strain IIIB) diluted

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to give a viral multiplicity of infection of 100 x TCID₅₀. A similar cell aliquot was diluted with medium to provide a mock-infected control. Cell infection was allowed to proceed for 1 hr at 37° in a tissue culture incubator with humidified 5% CO₂ atmosphere. After the 1 hr incubation the virus/cell suspensions were diluted 6-fold with fresh medium, and 125 µl of the cell suspension was added to each well of the plate containing pre-diluted compound. Plates were then placed in a tissue culture incubator with humidified 5% CO₂ for 5 days. At the end of the incubation period, 27 µl of 5% Nonidet-40 was added to each well of the incubation plate. After thorough mixing with a Costar multtipipetter, 60 µl of the mixture was transferred to filter-bottomed 96-well plates. The plates were analyzed in an automated assay instrument (Screen Machine, Idexx Laboratories). The assay makes use of a propidium iodide dye to estimate the DNA content of each well.

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1. Averett, D.R. 1989. Anti-HIV compound assessment by two novel high capacity assays. *J. Virol. Methods* **23**: 263-276.
2. Schwartz, O., et al. 1988. A rapid and simple colorimetric test for the study of anti-HIV agents. *AIDS Res. and Human Retroviruses*, **4**(6):441-447.
3. Daluge, S.M., et al. 1994. 5-chloro-2'3'-deoxy-3'fluorouridine (935U83), a selective anti-human immuno-deficiency virus agent with an improved metabolic and toxicological profile. *Antimicro. Agents and Chemother.*, **38** (7):1590-1603.

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The anti-viral potency in MT-4 cells of the compounds set forth in Tables 1 and 2 was determined using the above technique. The results are shown in Table 12.

5

Table 12.

<u>Cmpd#</u>	<u>K_i</u>	<u>IC₅₀</u>	<u>Cmpd#</u>	<u>K_i</u>	<u>IC₅₀</u>	<u>Cmpd#</u>	<u>K_i</u>	<u>IC₅₀</u>
1	A	G	123	C	NA	240	A	E
2	A	H	124	A	E	241	A	F
3	A	H	125	A	H	242	A	E
4	B	F	127	C	NA	243	A	E
5	A	H	201	B	H	244	A	F
6	A	NA	202	B	H	245	A	F
7	A	F	203	A	G	246	A	E
8	A	E	204	B	G	247	A	E
9	A	H	205	A	G	248	A	E
10	A	E	206	A	E	249	A	E
11	A	E	207	A	E	250	A	E
12	A	E	208	A	F	251	A	E
13	A	E	209	A	F	252	A	E
14	A	G	210	A	E	253	A	E
15	A	G	211	A	E	254	A	E
16	A	E	212	A	E	255	A	E
17	A	F	213	A	E	256	A	E
18	A	F	214	A	E	257	A	F
19	A	F	215	A	E	258	A	F
20	B	G	216	A	E	259	A	E
21	B	G	217	A	F	260	A	E
22	A	F	218	A	E	261	A	E
23	A	E	219	A	E	262	A	F
24	NA	NA	220	A	E	263	A	E
25	A	E	221	A	E	264	A	E

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<u>Compd#</u>	<u>Ki</u>	<u>IC50</u>	<u>Compd#</u>	<u>Ki</u>	<u>IC50</u>	<u>Compd#</u>	<u>Ki</u>	<u>IC50</u>
26	A	F	222	A	G	265	A	E
28	A	F	223	A	E	266	A	E
29	A	F	224	A	E	267	A	E
30	A	E	225	A	H	268	A	E
31	A	E	226	A	E	269	A	E
32	A	F	227	A	E	270	A	E
33	C	G	228	A	F	271	A	E
35	A	E	229	A	E	272	A	E
36	A	E	230	A	E	273	A	F
37	A	E	231	A	E	274	A	F
38	A	E	232	A	H	277	B	NA
39	A	E	233	A	F	278	B	NA
40	A	E	234	A	NA	279	C	NA
41	A	E	235	A	E	280	A	F
42	A	F	236	A	E	281	B	F
43	A	F	237	A	E			
44	A	NA	238	A	F			
113	C	G	239	A	E			

Biological Activity for Compounds 282-616

- Compounds 295-616 all had Ki<1nM (activity range "A") against the protease enzyme and IC50<100nM (activity range "E") in the antiviral assay unless specified differently below.
- Compounds 295-297, 402, 520, 523, 526, 527, 528, 540, 542, 563, 568, 605 had Ki in range "A" and IC50 in range "F".
- Compounds 529, 536, 543 had Ki in range "A" but IC50 not available.
- Compounds 506, 507, 508, 511, 560, 561, 564 had IC50 range >"E".

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4. Compounds 506, 507, 508, 511, 560, 561, 564 had IC₅₀ range >"E".

In Table 12 and Biological Activity for
Compounds 282-616, the following classifications
have been employed:

"I": K_i of less than 0.005 pM;

"A": K_i of less than 1 nM;

"B": K_i between 1 and 10 nM;

"C": K_i between 10 and 100 nM;

"D": K_i greater than 100 nM;

"J": IC₅₀ between 1.0 and 10.0 nM;

"E": IC₅₀ of 0.1 μM or less;

"F": IC₅₀ between 0.1 and 0.5 μM;

"G": IC₅₀ between 0.5 and 1.0 μM;

"H": IC₅₀ greater than 1.0 μM.

The designation "NA" is used where a given
compound was not tested.

Example

A comparison of K_i and IC₅₀ values for the
nitriles and amides of the present invention versus the
best compound, 4-amino-N-((2 syn, 3S)-2-hydroxy-4-phenyl-
3-((S)-tetrahydrofuran-3-ylloxycarbonylamino)-butyl)-N-
isobutyl-benzenesulfonamide (Compound A) disclosed in
International Publication W094/05639 is set forth in
Table 13.

4-Amino-N-((2 syn, 3S)-2-hydroxy-4-phenyl-3-
((S)-tetrahydrofuran-3-ylloxycarbonylamino)-butyl)-N-
isobutyl-benzenesulfonamide (Compound A) has a K_i of 75pM
against the enzyme and an IC₅₀ of 80 nM in the MT-4 cell
based assay.

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Table 13

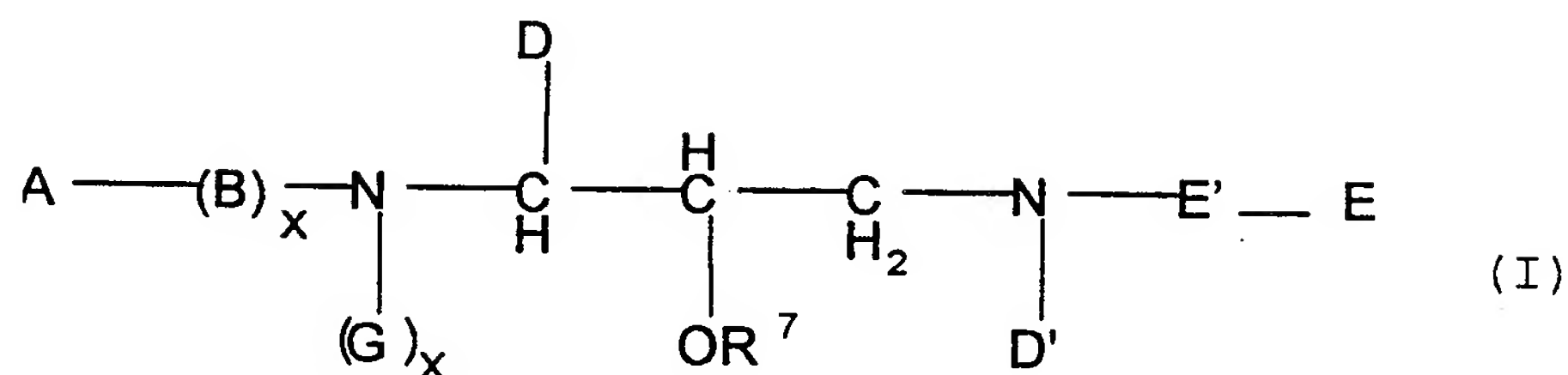
Compound	Ki (pM)	IC50 (nM)
12	I	J
39	I	J
374	I	J
375	I	J
376	I	J
377	I	J
378	I	J
379	I	J
A	75	80

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CLAIMS

We claim:

1. A compound of formula I:



or a pharmaceutically acceptable salt thereof, wherein:

E' is -CO- or -SO₂-;

A is selected from H; Ht; -R¹-Ht; -R¹-C₁-C₆ alkyl, which is optionally substituted with one or more groups independently selected from hydroxy, C₁-C₄ alkoxy, Ht, -O-Ht, -NR²-CO-N(R²)₂, -SO₂-R² or -CO-N(R²)₂; -R¹-C₂-C₆ alkenyl, which is optionally substituted with one or more groups independently selected from hydroxy, C₁-C₄ alkoxy, Ht, -O-Ht, -NR²-CO-N(R²)₂ or -CO-N(R²)₂; or R⁷;

each R¹ is independently selected from -C(O)-, -S(O)₂-, -C(O)-C(O)-, -O-C(O)-, -O-S(O)₂-, -NR²-S(O)₂-, -NR²-C(O)- or -NR²-C(O)-C(O)-;

each Ht is independently selected from C₃-C₇ cycloalkyl; C₅-C₇ cycloalkenyl; C₆-C₁₄ aryl; or a 5-7 membered saturated or unsaturated heterocycle, containing one or more heteroatoms selected from N, O, or S; wherein said aryl or said heterocycle is optionally fused to Q; and wherein any member of said Ht is optionally substituted with one or more substituents independently selected from oxo, -OR², SR², -R², -N(R²)(R²), -R²-OH, -CN,

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$-\text{CO}_2\text{R}^2$, $-\text{C}(\text{O})-\text{N}(\text{R}^2)_2$, $-\text{S}(\text{O})_2-\text{N}(\text{R}^2)_2$, $-\text{N}(\text{R}^2)-\text{C}(\text{O})-\text{R}^2$, $-\text{N}(\text{R}^2)-\text{C}(\text{O})\text{O}-\text{R}^2$, $-\text{C}(\text{O})-\text{R}^2$, $-\text{S}(\text{O})_n-\text{R}^2$, $-\text{OCF}_3$, $-\text{S}(\text{O})_n-\text{Q}$, methylenedioxy, $-\text{N}(\text{R}^2)-\text{S}(\text{O})_2(\text{R}^2)$, halo, $-\text{CF}_3$, $-\text{NO}_2$, Q , $-\text{OQ}$, $-\text{OR}^7$, $-\text{SR}^7$, $-\text{R}^7$, $-\text{N}(\text{R}^2)(\text{R}^7)$ or $-\text{N}(\text{R}^7)_2$;

each Q is independently selected from a 3-7 membered saturated, partially saturated or unsaturated carbocyclic ring system; or a 5-7 membered saturated, partially saturated or unsaturated heterocyclic ring containing one or more heteroatoms selected from O, N, or S; wherein Q is optionally substituted with one or more groups selected from oxo, $-\text{OR}^2$, $-\text{R}^2$, $-\text{SO}_2\text{R}^2$, $-\text{SO}_2-\text{N}(\text{R}^2)_2$, $-\text{N}(\text{R}^2)_2$, $-\text{N}(\text{R}^2)-\text{C}(\text{O})-\text{R}^2$, $-\text{R}^2-\text{OH}$, $-\text{CN}$, $-\text{CO}_2\text{R}^2$, $-\text{C}(\text{O})-\text{N}(\text{R}^2)_2$, halo, $-\text{CF}_3$;

each R^2 is independently selected from H, or C_1-C_4 alkyl,; and wherein said alkyl, when not a substituent of Q , is optionally substituted with Q or $-\text{OR}^3$; wherein when said R^2 is an $-\text{OR}^3$ substituted moiety, said R^3 in $-\text{OR}^3$ may not be $-\text{OR}^2$ substituted;

B , when present, is $-\text{N}(\text{R}^2)-\text{C}(\text{R}^3)_2-\text{C}(\text{O})-$;

each x is independently 0 or 1;

each R^3 is independently selected from H, Ht, C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_3-C_6 cycloalkyl or C_5-C_6 cycloalkenyl; wherein any member of said R^3 , except H, is optionally substituted with one or more substituents selected from $-\text{OR}^2$, $-\text{C}(\text{O})-\text{NH}-\text{R}^2$, $-\text{S}(\text{O})_n-\text{N}(\text{R}^2)(\text{R}^2)$, $-\text{N}(\text{R}^2)_2$, $-\text{N}(\text{R}^2)-\text{C}(\text{O})-\text{O}(\text{R}^2)$, $-\text{N}(\text{R}^2)-\text{C}(\text{O})-\text{N}(\text{R}^2)$, $-\text{N}(\text{R}^2)-\text{C}(\text{O})-(\text{R}^2)$, Ht, $-\text{CN}$, $-\text{SR}^2$, $-\text{CO}_2\text{R}^2$, or $\text{NR}^2-\text{C}(\text{O})-\text{R}^2$;

each n is independently 1 or 2;

G , when present, is selected from H, R^7 or C_1-C_4 alkyl, or, when G is C_1-C_4 alkyl, G and R^7 are optionally bound to one another either directly or through a C_1-C_3 linker to form a heterocyclic ring; or

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when G is not present, the nitrogen to which G is attached is bound directly to the R^7 group in $-OR^7$ with the concomitant displacement of one $-ZM$ group from R^7 ;

D is selected from Q; C_1 - C_6 alkyl optionally substituted with one or more groups selected from C_3 - C_6 cycloalkyl, $-OR^2$, $-S-Ht$, $-R^3$, $-O-Q$ or Q; C_2 - C_4 alkenyl optionally substituted with one or more groups selected from $-OR^2$, $-S-Ht$, $-R^3$, $-O-Q$ or Q; C_3 - C_6 cycloalkyl optionally substituted with or fused to Q; or C_5 - C_6 cycloalkenyl optionally substituted with or fused to Q;

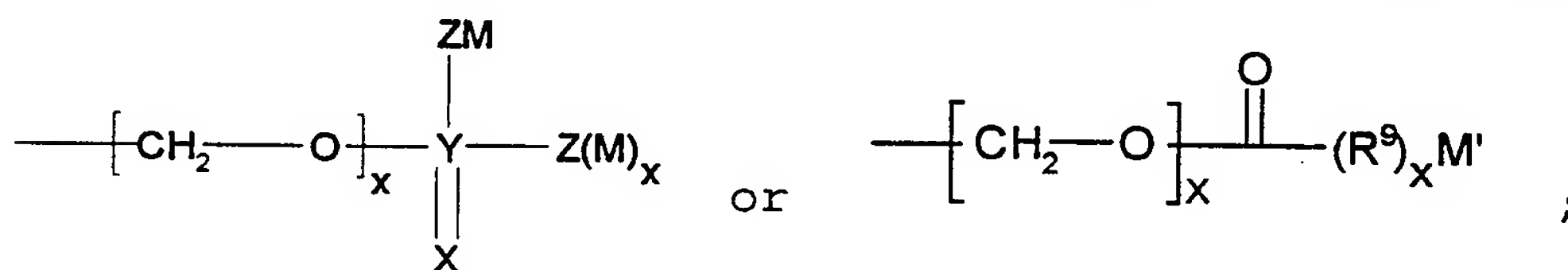
D' is selected from C_1 - C_{15} alkyl, C_2 - C_{15} alkenyl or C_2 - C_{15} alkynyl, each of which contains one or more substituents selected from oxo, halo, $-CF_3$, $-OCF_3$, $-NO_2$, azido, $-SH$, $-SR^3$, $-N(R^3)-N(R^3)_2$, $-O-N(R^3)_2$, $-(R^3)N-O-(R^3)$, $-N(R^3)_2$, $-CN$, $-CO_2R^3$, $-C(O)-N(R^3)_2$, $-S(O)_n-N(R^3)_2$, $-N(R^3)-C(O)-R^3$, $-N(R^3)-C(O)-N(R^3)_2$, $-N(R^3)-C(O)-S(R^3)$, $-C(O)-R^3$, $-S(O)_n-R^3$, $-N(R^3)-S(O)_n(R^3)$, $-N(R^3)-S(O)_n-N(R^3)_2$, $-S-NR^3-C(O)R^3$, $-C(S)N(R^3)_2$, $-C(S)R^3$, $-NR^3-C(O)OR^3$, $-O-C(O)OR^3$, $-O-C(O)N(R^3)_2$, $-NR^3-C(S)R^3$, $=N-OH$, $=N-OR^3$, $=N-N(R^3)_2$, $=NR^3$, $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$, $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-OR^3$, $-N(R^3)-C[=N-CN]-OR^3$, $-N(R^3)-C[=N-CN]-(R^3)_2$, $-OC(O)R^3$, $-OC(S)R^3$, $-OC(O)N(R^3)_2$, $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-N(R^3)_2$, $O-C(O)N(OR^3)(R^3)$, $N(R^3)-N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $-OC(S)N(R^3)_2$, $-OC(S)N(R^3)(R^3)$, or PO_3-R^3 ; with the proviso that when R^7 is H, E' is $-SO_2$, G is H or alkyl, and when B is present or when B is not present and R^1 is $-C(O)-$, D' may not be C_1 - C_{15} alkyl substituted with one substituent selected from $-N(R^3)_2$, $-SR^3$ or $-S(O)_n-R^3$, or substituted with two $-N(R^3)_2$ substituents;

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E is selected from Ht; O-Ht; Ht-Ht; Ht fused with Ht; $-O-R^3$; $-N(R^2)(R^3)$; C_1-C_6 alkyl optionally substituted with one or more groups selected from R^4 or Ht; C_2-C_6 alkenyl optionally substituted with one or more groups selected from R^4 or Ht; C_3-C_6 saturated carbocycle optionally substituted with one or more groups selected from R^4 or Ht; or C_5-C_6 unsaturated carbocycle optionally substituted with one or more groups selected from R^4 or Ht;

each R^4 is independently selected from $-OR^2$, $-OR^3$, $-SR^2$, $-SOR^2$, $-SO_2R^2$, $-CO_2R^2$, $-C(O)-NHR^2$, $-C(O)-N(R^2)_2$, $-C(O)-NR^2(OR^2)$, $-S(O)_2-NHR^2$, halo, $-NR^2-C(O)-R^2$, $-N(R^2)_2$ or $-CN$;

each R^7 is independently selected from hydrogen,



wherein each M is independently selected from H, Li, Na, K, Mg, Ca, Ba, $-N(R^2)_4$, C_1-C_{12} -alkyl, C_2-C_{12} -alkenyl, or $-R^6$; wherein 1 to 4 $-CH_2$ radicals of the alkyl or alkenyl group, other than the $-CH_2$ that is bound to Z, is optionally replaced by a heteroatom group selected from O, $S(O)$, $S(O)_2$, or $N(R^2)$; and wherein any hydrogen in said alkyl, alkenyl or R^6 is optionally replaced with a substituent selected from oxo, $-OR^2$, $-R^2$, $N(R^2)_2$, $N(R^2)_3$, R^2OH , $-CN$, $-CO_2R^2$, $-C(O)-N(R^2)_2$, $S(O)_2-N(R^2)_2$, $N(R^2)-C(O)-R^2$, $C(O)R^2$, $-S(O)_n-R^2$, OCF_3 , $-S(O)_n-R^6$, $N(R^2)-S(O)_2(R^2)$, halo, $-CF_3$, or $-NO_2$;

M' is H, C_1-C_{12} -alkyl, C_2-C_{12} -alkenyl, or $-R^6$; wherein 1 to 4 $-CH_2$ radicals of the alkyl or alkenyl group is

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optionally replaced by a heteroatom group selected from O, S, S(O), S(O)₂, or N(R²); and wherein any hydrogen in said alkyl, alkenyl or R⁶ is optionally replaced with a substituent selected from oxo, -OR², -R², -N(R²)₂, N(R²)₃, -R²OH, -CN, -CO₂R², -C(O)-N(R²)₂, -S(O)₂-N(R²)₂, -N(R²)-C(O)-R₂, -C(O)R², -S(O)_n-R², -OCF₃, -S(O)_n-R⁶, -N(R²)-S(O)₂(R²), halo, -CF₃, or -NO₂;

Z is O, S, N(R²)₂, or, when M is not present, H.

Y is P or S;

X is O or S;

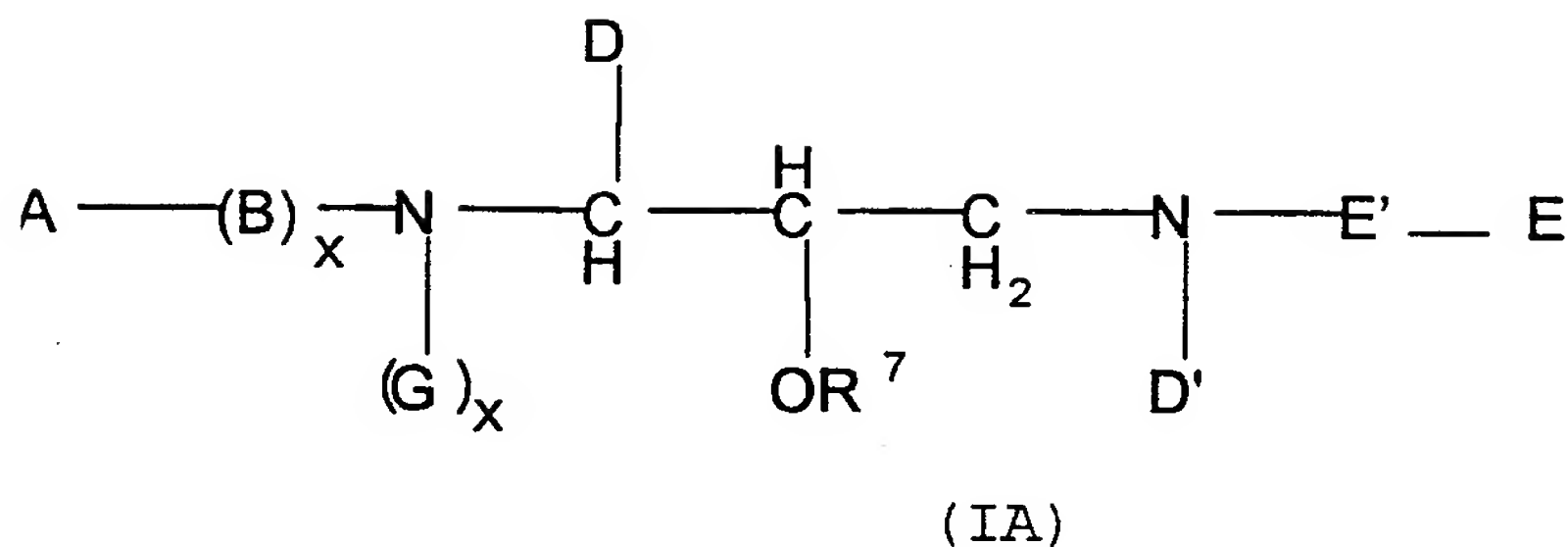
R⁹ is C(R²)₂, O or N(R²); and wherein when Y is S, Z is not S;

R⁶ is a 5-6 membered saturated, partially saturated or unsaturated carbocyclic or heterocyclic ring system, or an 8-10 membered saturated, partially saturated or unsaturated bicyclic ring system; wherein any of said heterocyclic ring systems contains one or more heteroatoms selected from O, N, S, S(O)_n or N(R²); and wherein any of said ring systems optionally contains 1 to 4 substituents independently selected from OH, C₁-C₄ alkyl, -O-C₁-C₄ alkyl or -O-C(O)-C₁-C₄ alkyl; and

each R⁵ is independently selected from hydrogen, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl or Ht, wherein any R⁵, except for hydrogen, is optionally substituted with -CF₃, -PO₃R³, azido or halo.

2. The compound according to claim 1, having the formula IA:

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wherein:

D' is selected from C₁₋₁₅ alkyl, C₂₋₁₅ alkenyl or C₂₋₁₅ alkynyl; each of which is substituted with one to two -CN groups and each of which is optionally substituted with C₃₋₈ cycloalkyl.

3. The compound according to claim 2

wherein:

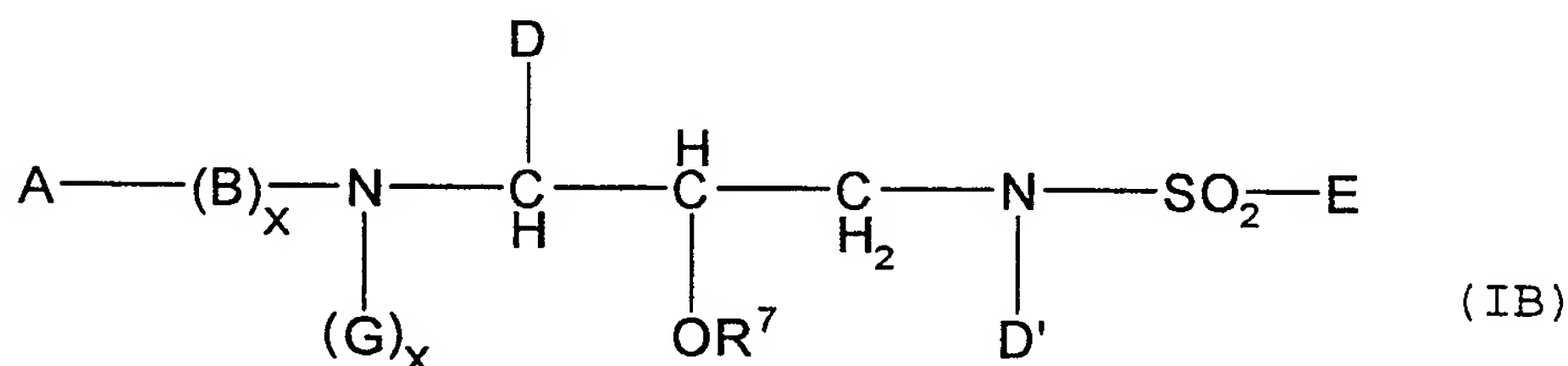
D' is selected from C₁₋₁₅ alkyl or C₂₋₁₅ alkenyl; each of which is substituted with one to two -CN groups and each of which is optionally substituted with C₃₋₈ cycloalkyl.

4. The compound according to claim 2

wherein:

D' is C₂₋₁₅ alkynyl which is substituted with one to two -CN groups and each of which is optionally substituted with C₃₋₈ cycloalkyl.

5. The compound according to claim 1 having the formula IB:



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wherein:

D' is selected from C₁-C₁₅ alkyl, C₂-C₁₅ alkenyl or C₂-C₁₅ alkynyl, each of which contains one or more substituents selected from oxo, halo, -CF₃, -OCF₃, -NO₂, azido, -SH, -SR³, -N(R³)-N(R³)₂, -O-N(R³)₂, -(R³)N-O-(R³), -N(R³)₂, -CO₂R³, -C(O)-N(R³)₂, -S(O)_n-N(R³)₂, -N(R³)-C(O)-R³, -N(R³)-C(O)-N(R³)₂, -N(R³)-C(O)-S(R³), -C(O)-R³, -S(O)_n-R³, -N(R³)-S(O)_n(R³), -N(R³)-S(O)_n-N(R³)₂, -S-NR³-C(O)R³, -C(S)N(R³)₂, -C(S)R³, -NR³-C(O)OR³, -O-C(O)OR³, -O-C(O)N(R³)₂, -NR³-C(S)R³, =N-OH, =N-OR³, =N-N(R³)₂, =NR³, =NNR³-C(O)N(R³)₂, =NNR³-C(O)OR³, =NNR³-S(O)_n-N(R³)₂, -NR³-C(S)OR³, -NR³-C(S)N(R³)₂, -NR³-C[=N(R³)]-N(R³)₂, -N(R³)-C[=N-NO₂]-N(R³)₂, -N(R³)-C[=N-NO₂]-OR³, -N(R³)-C[=N-CN]-OR³, -N(R³)-C[=N-CN]-(R³)₂, -OC(O)R³, -OC(S)R³, -OC(O)N(R³)₂, -C(O)N(R³)-N(R³)₂, -O-C(O)N(R³)-N(R³)₂, O-C(O)N(OR³)(R³), N(R³)-N(R³)-C(O)R³, N(R³)-OC(O)R³, N(R³)-OC(O)R³, N(R³)-OC(O)R³, -OC(S)N(R³)₂, -OC(S)N(R³)(R³), or PO₃-R³; with the proviso that when R⁷ is H, E' is -SO₂-, G is H or alkyl, and when B is present or when B is not present and R¹ is -C(O)-, D' may not be C₁-C₁₅ alkyl substituted with one substituent selected from -N(R³)₂, -SR³ or -S(O)_n-R³, or substituted with two -N(R³)₂ substituents.

6. The compound according to claim 5 wherein:

D' is selected from C₁-C₁₅ alkyl or C₂-C₁₅ alkenyl, each of which contains one or more substituents selected from oxo, halo, -CF₃, -OCF₃, -NO₂, azido, -N(R³)-N(R³)₂, -O-N(R³)₂, -(R³)N-O-(R³), -N(R³)₂, -N(R³)-C(O)-N(R³)₂, -N(R³)-C(O)-S(R³), -C(O)-R³, -S(O)_n-R³, -N(R³)-S(O)_n(R³), -N(R³)-

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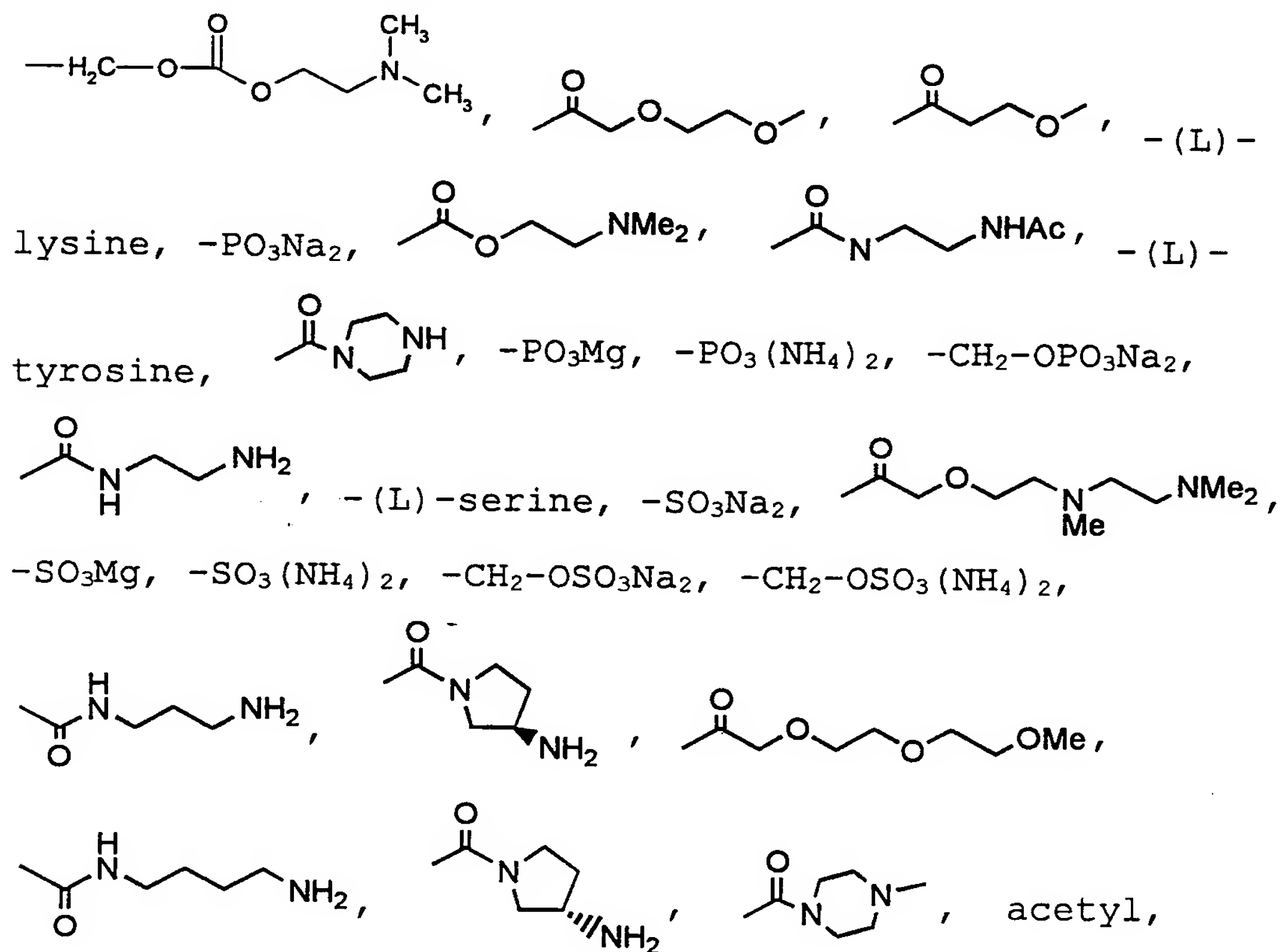
$S(O)_n-N(R^3)_2$, $-S-NR^3-C(O)R^3$, $-C(S)N(R^3)_2$, $-C(S)R^3$, $-NR^3-C(O)OR^3$, $-O-C(O)OR^3$, $-O-C(O)N(R^3)_2$, $-NR^3-C(S)R^3$, $=N-OH$, $=N-OR^3$, $=N-N(R^3)_2$, $=NR^3$, $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$, $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-OR^3$, $-N(R^3)-C[=N-CN]-OR^3$, $-N(R^3)-C[=N-CN]-(R^3)_2$, $-OC(O)R^3$, $-OC(S)R^3$, $-OC(O)N(R^3)_2$, $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-N(R^3)_2$, $O-C(O)N(OR^3)(R^3)$, $N(R^3)-N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $-OC(S)N(R^3)_2$, $-OC(S)N(R^3)(R^3)$, or PO_3-R^3 ; C_2-C_{15} alkynyl which contains one or more substituents selected from oxo, halo, $-CF_3$, $-OCF_3$, $-NO_2$, azido, $-SH$, $-SR^3$, $-N(R^3)-N(R^3)_2$, $-O-N(R^3)_2$, $-(R^3)N-O-(R^3)$, $-N(R^3)_2$, $-CO_2R^3$, $-C(O)-N(R^3)_2$, $-S(O)_n-N(R^3)_2$, $-N(R^3)-C(O)-R^3$, $-N(R^3)-C(O)-N(R^3)_2$, $-N(R^3)-C(O)-S(R^3)$, $-C(O)-R^3$, $-S(O)_n-R^3$, $-N(R^3)-S(O)_n(R^3)$, $-N(R^3)-S(O)_n-N(R^3)_2$, $-S-NR^3-C(O)R^3$, $-C(S)N(R^3)_2$, $-C(S)R^3$, $-NR^3-C(O)OR^3$, $-O-C(O)OR^3$, $-O-C(O)N(R^3)_2$, $-NR^3-C(S)R^3$, $=N-OH$, $=N-OR^3$, $=N-N(R^3)_2$, $=NR^3$, $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$, $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-OR^3$, $-N(R^3)-C[=N-CN]-OR^3$, $-N(R^3)-C[=N-CN]-(R^3)_2$, $-OC(O)R^3$, $-OC(S)R^3$, $-OC(O)N(R^3)_2$, $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-N(R^3)_2$, $O-C(O)N(OR^3)(R^3)$, $N(R^3)-N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $-OC(S)N(R^3)_2$, $-OC(S)N(R^3)(R^3)$, or PO_3-R^3 ; with the proviso that when R^7 is H, E' is $-SO_2-$, G is H or alkyl, and when B is present or when B is not present and R^1 is $-C(O)-$, D' may not be C_1-C_{15} alkyl substituted with one substituent selected from $-N(R^3)_2$ or $-S(O)_n-R^3$, or substituted with two $-N(R^3)_2$ substituents.

7. The compound according to claim 5 wherein:

D' is selected from C₁-C₁₅ alkyl or C₂-C₁₅ alkenyl, each of which contains one or more substituents selected from -SH, -SR³, -CO₂R³, -C(O)-N(R³)₂, -S(O)_n-N(R³)₂ or -N(R³)-C(O)-R³; with the proviso that when R⁷ is H, E' is -SO₂-, G is H or alkyl, and when B is present or when B is not present and R¹ is -C(O)-, D' may not be C₁-C₁₅ alkyl substituted with one substituent selected from -SR³.

8. The compound according to any one of claims 1 to 4, wherein E' is SO₂.

9. The compound according to any one of claims 1 to 7, wherein at least one R⁷ is selected from:

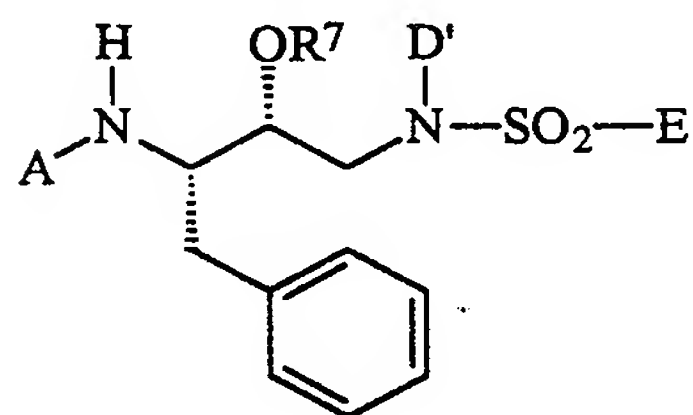


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CC(=O)C, CCCC(=O)C, -(L)-valine, -(L)-glutamic acid, -(L)-
aspartic acid, -(L)-γ-t-butyl-aspartic acid, CC(=O)OC1CCCOC1,
-(L)-(L)-3-pyridylalanine, -(L)-histidine, -CHO, CC(=O)CF3,
CC(=O)OC1CCCOC1, CC(=O)OCCCCCO[C@H]1[C@@H](OC(=O)C)[C@H](OC(=O)C)[C@H](OC(=O)C)[C@H]1OC(=O)C, [O-]P(=O)([O-])OCC[NH3+],
[O-]P(=O)([O-])OCCN(C)C, CCOP(=O)([O-])[O-], CCOS(=O)(=O)[O-], PO₃K₂, PO₃Ca,

PO₃-spermine, PO₃-(spermidine)₂ or PO₃-(meglamine)₂.

10. The compound according to claim 8, having the formula II:

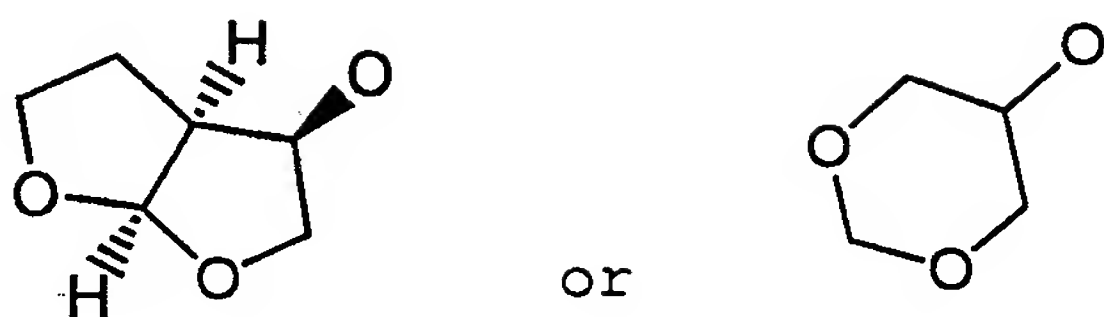


(II) .

11. The compound according to claim 10,
wherein:

A is R'-C(O)-; and
R' is selected from

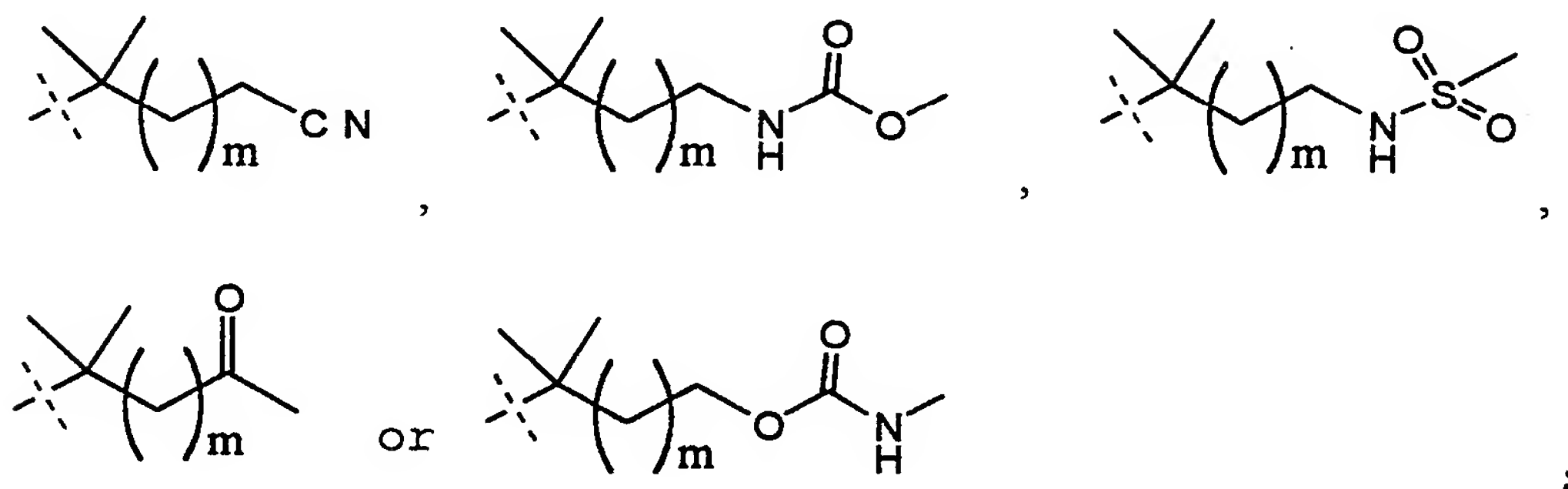
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12. The compound according to claim 10,
wherein:

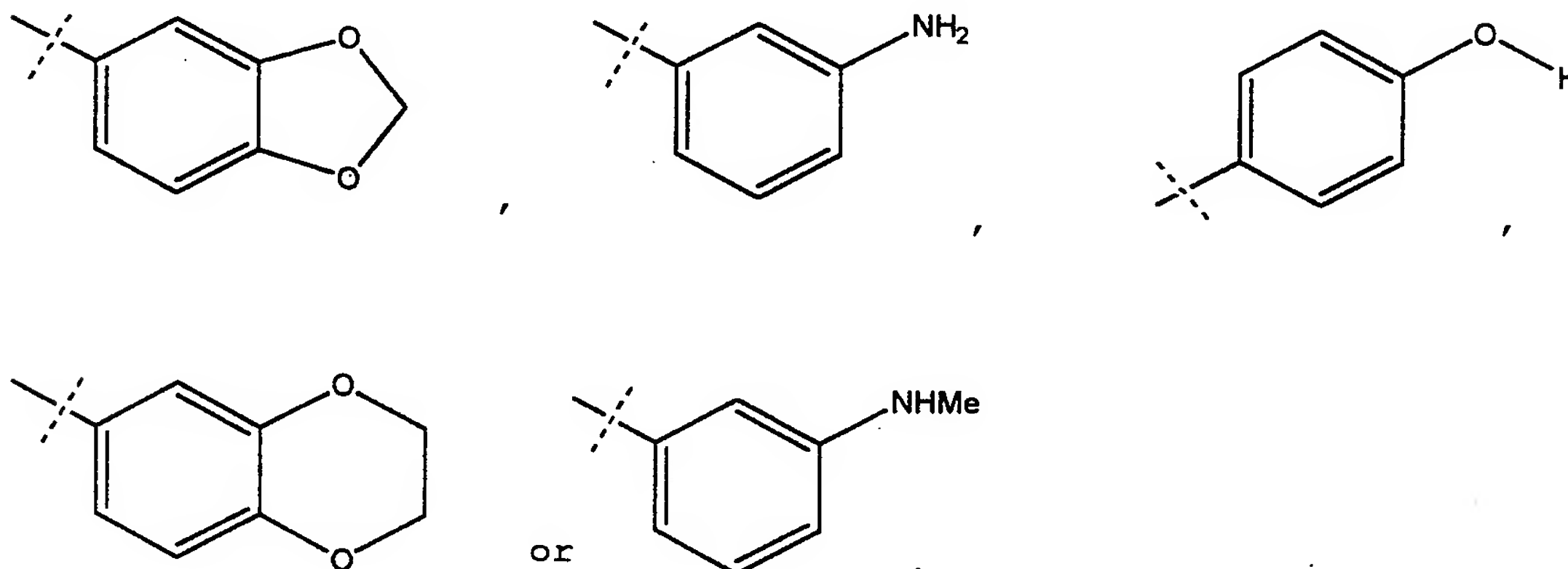
D' is $-\text{CH}_2-\text{R}''$; and

R'' is selected from



wherein m is 0 to 3.

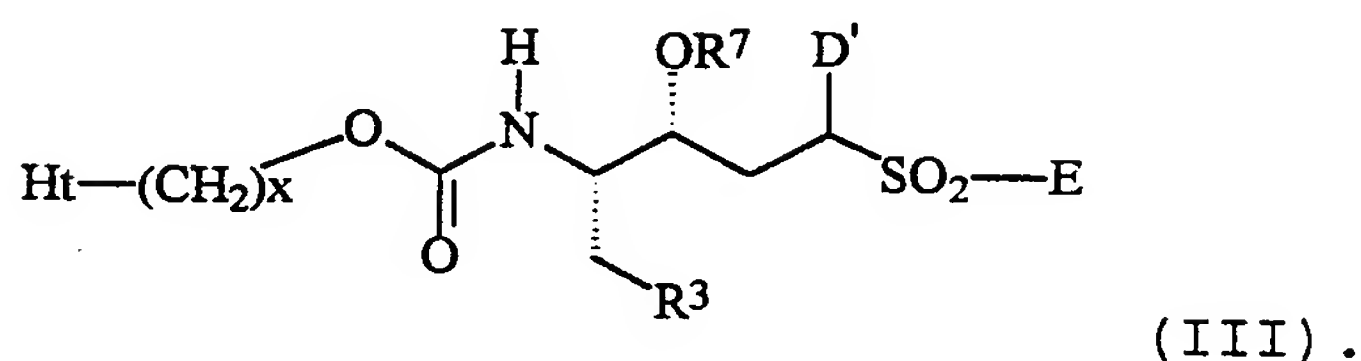
13. The compound according to claim 10,
wherein E is selected from



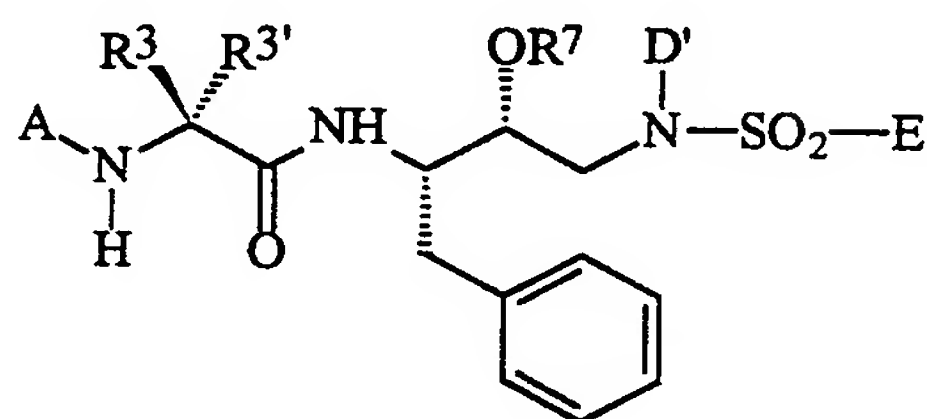
14. The compound according to claim 10,
wherein R⁷ is $-\text{PO}_3^{2-}$.

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15. The compound according to claim 1, having the formula III:



16. The compound according to claim 1, having the formula IV:



wherein $R^{3'}$ is selected from H, Ht, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_3 - C_6 cycloalkyl or C_5 - C_6 cycloalkenyl; wherein any member of said R^3 , except H, is optionally substituted with one or more substituents selected from $-OR^2$, $-C(O)-NH-R^2$, $-S(O)_n-N(R^2)(R^2)$, $-N(R^2)_2$, $-N(R^2)-C(O)-O(R^2)$, $-N(R^2)-C(O)-N(R^2)$, $-N(R^2)-C(O)-(R^2)$, $-N(R^2-OR^2)_2$, $-C(O)-Ht$, Ht, $-CN$, $-SR^2$, $-CO_2R^2$, or $NR^2-C(O)-R^2$.

17. The compound according to claim 11, wherein said compound is selected from any one of compound numbers: 210, 224, 240, 248, 250, 255, 263, 270, 272, 280, 299, 300, 307, 309, 313, 314, 315, 316, 359, 360, 384, 483, 494, 496, 523, 524, 531, 542, 548, 553, 558, 563, 570, 571, 575, 579, 589, 596, 606, 609, 616.

18. The compound according to claim 11, wherein said compound is selected from any one of

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compound numbers: 12, 16, 25, 29, 30, 31, 35, 39, 41, 42, 47, 100, 124, 375, 378, 421, 459, 464.

19. The compound according to claim 17, wherein said compound is selected from any one of compound numbers: 224, 240, 263, 270, 272, 280, 299, 300, 307, 309, 313, 314, 315, 316, 359, 360, 384, 483, 494, 496, 548, 553, 558, 563, 570, 571, 575, 579, 589, 596, 606, 609, 616.

20. The compound according to claim 18, wherein said compound is selected from any one of compound numbers: 12, 16, 25, 35, 39, 42, 47, 100, 375, 378, 421, 459, 464.

21. The compound according to claim 19, wherein said compound is selected from any one of compound numbers: 224, 240, 272, 299, 314.

22. The compound according to claim 20, wherein said compound is selected from any one of compound numbers: 16, 25, 42, 47, 100.

23. A composition comprising a compound according to any one of claims 1-22 or a pharmaceutically acceptable salt thereof in an amount sufficient to detectably inhibit aspartyl protease activity in a patient, and a pharmaceutically acceptable carrier.

24. The composition according to claim 23, further comprising an additional antiviral agent other than a compound of formula (I).

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25. The composition according to claim 23, wherein said composition is formulated as a pharmaceutically acceptable, orally available tablet or capsule.

26. A method of treating an HIV virus infection in a human comprising the step of administering to said human a composition according to any one of claims 23 to 25.

27. The method according to claim 26, comprising the step of administering to said patient an additional antiviral agent other than a compound of formula I, wherein said additional antiviral agent is administered prior to, simultaneously with or following administration of said composition.

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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: INHIBITORS OF ASPARTYL PROTEASE

(57) Abstract: The present invention relates to a novel class of sulfonamides which are aspartyl protease inhibitors. In one embodiment, this invention relates to a novel class of HIV aspartyl protease inhibitors characterized by specific structural and physicochemical features. This invention also relates to pharmaceutical compositions comprising these compounds. The compounds and pharmaceutical compositions of this invention are particularly well suited for inhibiting HIV-1 and HIV-2 protease activity and consequently may be advantageously used as anti-viral agents against the HIV-1 and HIV-2 viruses. This invention also relates to methods for inhibiting the activity of HIV aspartyl protease using the compounds of this invention and methods for screening compounds for anti-HIV activity.



WO 00/47551 A3

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 00/03288

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D493/04 C07C311/29 C07C311/42 C07D317/62 C07D307/20
C07D319/06 C07D407/12 C07F9/6561 A61K31/34 A61K31/36
A61K31/18 A61K31/665

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D C07C C07F A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 94 04492 A (G.D. SEARLE & CO ET AL) 3 March 1994 (1994-03-03) page 40, lines 26-29; table 1A, entries 34, 37-38; table 16, entry 10; page 157, lines 21-34	1,5-8, 10,15, 23-27
A	page 3, line 21- page 5, line 34; table 4, entries 81, 84-85; page 157, lines 21-34	1,5-8, 16,23-27
X	WO 94 10136 A (G.D. SEARLE & CO ET AL) 11 May 1994 (1994-05-11) table 1c, entry J	1,5,7,8, 10,15
A	page 3, line 9 - page 5, line 26; table 3, entries 64, 67, 68; page 87, line 28 - page 88, line 4	1,5-8, 10,23-27
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☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/03288

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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A	A.K. GHOSH ET AL: BIOORG. MED. CHEM. LETT., vol. 8, no. 6, 1998, pages 687-690, XP004136945 table 1 ---	1,8,10, 11,23,26
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Information on patent family members

International Application No

PCT/US 00/03288

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